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(U) SRI INTERNATIONAL MENLO PARK CA LIFE SCIENCES DIV  
C S REBERT ET AL. JUN 83 AFOSR-TR-83-0902  
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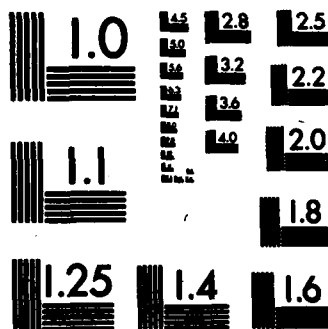
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## NEUROPHYSIOLOGICAL BASES OF EVENT-RELATED POTENTIALS

Annual Report No. 1

June 1983

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By: Charles S. Rebert and William J. Donovan  
SRI International

and

Karl H. Pribram and Jeffrey E. Evans  
Stanford University

Prepared for:

AIR FORCE OFFICE OF SCIENTIFIC RESEARCH  
Life Sciences Directorate  
Bolling AFB, D.C. 20332

Attention: Dr. Alfred R. Fregly  
Program Manager

AFOSR Contract No. F49620-82-K-0016

SRI Project LSU 4373

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# SRI International



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20. ABSTRACT (Continue on reverse side if necessary and identify by block number) ✓ In order to more fully understand the physiological and psychological significance of event-related potentials, cortical and subcortical recordings are being obtained from monkeys performing in operant-conditioning tasks. Six animals were trained on initial phases of the cued reaction-time task at SRI International and were subsequently implanted with electrodes capable of recording transient and sustained evoked potentials and massed-unit activity. Two monkeys were trained on initial phases of an "oddball" task at Stanford University, and (Continued on reverse)			

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ABSTRACT (Concluded)

→ electrodes are being prepared so that the subcortical generators of the P300 wave can be assessed in these animals. An LSI-11/23 computer system was installed at SRI to implement the cued reaction-time task and to collect event-related potentials. Preliminary recordings of slow potentials and massed-unit activity were collected from the lateral geniculate nucleus of one cat to evaluate the performance of modified amplifiers, and transient (P300) and sustained (contingent negative variation) evoked potentials were recorded from the scalps of human subjects to confirm appropriate performance of the laboratory system. ↑

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## **A. Background and Purposes**

### **1. Man-Machine Systems**

The modern fighter pilot is primarily an "executive"--an information-processing and decision-making element of a complex man-machine system. The almost overwhelming amount of information to be processed from displays related to flight-systems status, navigation, communications, weapons-threat warning, radar, imaging sensor systems, and situational displays has precipitated the need for improvements in the engineering of cockpit displays and in the understanding of the human operator's information-processing characteristics (Reising, 1980; Furness, 1980) and workload parameters (Moray, 1978). Although the human operator is generally regarded as the weakest link in man-machine systems, the human element is critical if systems are to retain the capability of reacting intelligently and imaginatively to unanticipated conditions (Gomer et al., 1979).

Because pilot workload is now primarily "mental," the concepts and procedures of cognitive psychology are particularly relevant to the solution of workload problems and man-machine interfacing. Cognitive psychology has undergone a striking revolution within the last quarter century, involving greater emphasis on concepts such as information-processing (Simon, 1980) and intention (Jung, 1981; O'Connor, 1981). Early behaviorists generally considered "cognitions" such as thoughts, feelings, evaluations, and expectancies as epiphenomena that had no relevance to the mechanics of actual behavior, which was conceived to flow from particular stimulus events. However, as recently emphasized by O'Connor (1981), Jung (1981), and Donchin (1980), intentions and goals precede and precipitate (rather than result from) perceptual, attentional, and behavioral strategies.

Although the information-processing revolution has led to a synthesis of several dimensions of psychological research, there remains

a large gap in explanations of cognition in that little is known about its neural substrates. A complete understanding of human thinking will probably not be possible until the neural processes underlying symbol manipulations can be specified (Simon, 1980). Obviously, the more complete our knowledge of cognitive processes, the more thorough will be the solution of problems relating to the efficiency of man-machine systems.

## 2. Event-Related Brain Potentials

The only direct indications of brain function routinely available to the psychophysiolgist are electric fields accompanying "spontaneous" and event-related intracerebral activity. The slow-wave (1-20 Hz) electroencephalogram (EEG) provides a very general index of the patterning of "activation" across the cerebral mantle. Such measures can be useful in assessing the extent to which various cortical regions--for example, the left and right hemispheres--are differentially involved in various types of tasks (Rebert, 1980a).

Event-related potentials (ERPs) are patterns of electric change associated with the occurrence of fairly discrete external or internal events--a flash of light, a decision. Various components of ERPs reflect activity in different regions of the brain and different information-processing functions, but--with few exceptions--the exact source of the potentials and their precise relationships to cognition, effort, motivation, and overt behavior are unknown. These potentials range from the very specific click-evoked, high-frequency burst of waves generated in brainstem auditory structures (volume-conducted to scalp electrodes) to long-lasting DC potentials of the cortex related to anticipatory processes. Although ERPs are composite reflections of a myriad of intracerebral transactions and their true form is distorted by tissues between the cortex and scalp recording electrodes, they are extremely useful tools for assessing the functional integrity of the nervous system (Regan, 1972; Aminoff, 1980; Rebert, 1980b). ERPs have been the focus of interest of many psychophysiolgists interested in the neural correlates of cognitive processes (e.g., Donchin and Lindsay,

1969; Kornhuber and Deecke, 1980). Picton and Stuss (1980) have thoroughly summarized the component structure of the known ERPs, their sensitivities to various types of experimental manipulations, and their presumed relationships to psychological processes. The component structure of ERPs varies as a function of stimulus modality, recording location, task parameters, and subject state variables, among many other factors. In a situation requiring the detection of a rare event, a prominent positive wave (P300) occurs, with latency of about 300 msec. This may represent the response to disconfirmation of expectancy and is influenced by other subjective factors such as decision confidence (Hillyard et al., 1978).

In the cued reaction-time (RT) task, one stimulus acts as a warning that a second stimulus, which has significance for the subject, will subsequently appear. During the few seconds of the interstimulus interval, there appears a slow negative potential shift, called the contingent negative variation (CNV). This event is probably a nonspecific sign of localized cortical activation (Rebert, 1980c). A slow potential shift, the Bereitschaftspotential (BP), which is morphologically similar to the late portion of the CNV, occurs when a S prepares, in the absence of any preparatory or imperative cues, to carry out a behavioral act.

### 3. Basic Research in Animals

Although studies of human electrocortical activities demonstrate the validity of the "biocybernetic" concept (Donchin, 1980; Rebert, 1980a), a complete knowledge of ERPs using just those procedures is precluded by a number of limitations in human scalp-recording methods. For example, scalp recordings provide an extremely limited view of brain activity (i.e., scalp potentials are not precise reflections of the underlying cerebral activity because of distortions produced by intervening tissues); many cortical events are not apparent in scalp recordings; and ERP components recorded from the scalp are unlikely to be due to discrete generators, but probably reflect overlapping sources of potentials.

The foregoing considerations point clearly to the need for studies of ERPs in animals. The advantages of using animal subjects lie, of course, in the wide variety of procedures and experimental manipulations that can be carried out--e.g., intracerebral recording and stimulating (either electrically or pharmacologically), disruption of known neural pathways, histological evaluations, long-term study of a subject, systemic injection of a variety of pharmacological agents, direct manipulation of biological drive states by deprivation, and rigorous control over the experimental experiences of the subjects.

#### 4. Choice of Experimental Paradigm

A host of experimental paradigms can be employed with animals to study ERPs. The one selected should cognitively engage the animal and closely approximate paradigms used in human research. Most preferred is a paradigm that is sufficiently general to include a variety of psychological processes and ERP components, is rigorous in terms of good control over the behavioral sequences and psychological sets induced in the animal, and is flexible in terms of the ability to manipulate a variety of experimental variables while not altering the basic logical structure of the task. In addition, because homology between animal and human ERPs is important, advantages should accrue from the use of a behavioral paradigm for which there already exist data indicating a close homology of ERPs elicited by the situation (Rebert, 1972).

The cued RT task meets the foregoing criteria and was considered to be the most promising one to use in early studies of the electrogenesis of ERPs in animals.

Another major paradigm of interest in this research was the "oddball" task wherein rare and common events elicit different ERPs, the former evoking the P300 component, which appears to follow disconfirmation of expectancy and which might be involved in mnemonic processes (Donchin, 1979). To facilitate study of this process, funds were made available to Dr. Karl Pribram at Stanford University to begin experiments using visual stimuli to which monkeys make behavioral responses. The paradigms, equipment, and preliminary behavioral training at

Stanford are described later in this report. In addition, a passive P300 paradigm (not requiring behavioral responses from the monkeys) using tone stimuli has been implemented at SRI.

## 5. General Purposes

The broad goal of the proposed research is to describe the dynamics of the general cerebral system that is operative in the cued RT task, using as a primary tool the recording of ERPs and associated neuronal activity in various regions of the monkey's brain. Thus, the intra-cerebral distribution of ERPs, their behavioral significance and similarity to human ERPs, their relationship to neuronal activity, and mediation by chemical transmitter/modulator substances are to be studied. Later stages of the research will be directed toward discovering some of the anatomical/neurochemical pathways by which ERPs become modulated during the cued RT task.

This research will provide information that is necessary to a fuller understanding of ERPs, and thus will contribute to a more substantial understanding of brain processes involved in a variety of cognitive acts. Such an understanding should contribute to a more solid foundation for using ERPs in a biocybernetic system.

## B. Preparation of Facilities and Equipment

### 1. General Stages of Preparation

The SRI laboratory facility used for this research was developed in two stages. First, equipment previously used for similar purposes, including an Iconix Logic unit and Linc-8 computers, was configured so that preliminary training of six monkeys could be undertaken early in the project period. Subsequently, a DEC LSI-11/23 computer system, purchased by SRI for use on this project, was installed, and software and hardware elements were configured to provide more automatic and comprehensive experimental control, data acquisition, and analysis than that provided by the older equipment.

## 2. Iconix Logic Unit for Preliminary Training

A schematic representation of the logic of the cued RT task is shown in Figure 1. A trial can be initiated if the animal has maintained a specified hand posture (holding on to a round knob attached to the primate chair) for at least 5 sec. After a period of training, the position is usually maintained throughout the intertrial interval (ITI). This contingency assures a greater homogeneity of RT because the instrumental response is always made from the same starting position. Tone bursts (1 or 3 kHz), 100-msec and 72 dB (re: 20  $\mu$ N/m<sup>2</sup>), constitute warning or discriminative stimuli (WS and DS, respectively). The WS is followed by an imperative stimulus (IS), a light, which indicates to the monkey that it can obtain reinforcement by making the appropriate operant response (a bar press in this case). The interstimulus interval is typically 1.5 sec, but can be manipulated for experimental reasons. If the monkey releases the "hold" position any time before onset of the IS, the trial is aborted and no reward is available. Correct performance allows the monkey to receive 1 ml of an orange-flavored drink (Tang®) for each bar-press made during the 12 sec that the IS remains on (usually a total of 15-20 ml).

The DS occurs in isolation--i.e., it is not paired with any other cue--and provides a comparison for assessing ERP components related to the associative responses elicited by the WS. Typically, CNVs are evoked by both the WS and DS early in the training period, but later only by the WS. Thus, this paradigm permits assessment of the development of associative and discriminative events in several regions of the brain (Rebert, 1977).

The many contingencies in this paradigm required an elaborate and fairly time-consuming programming of the Iconix unit. For example, trials are not initiated unless the animal has sustained a fixed position of the right hand for at least 5 sec, the availability of reinforcement is contingent on the presence of the IS, trials are



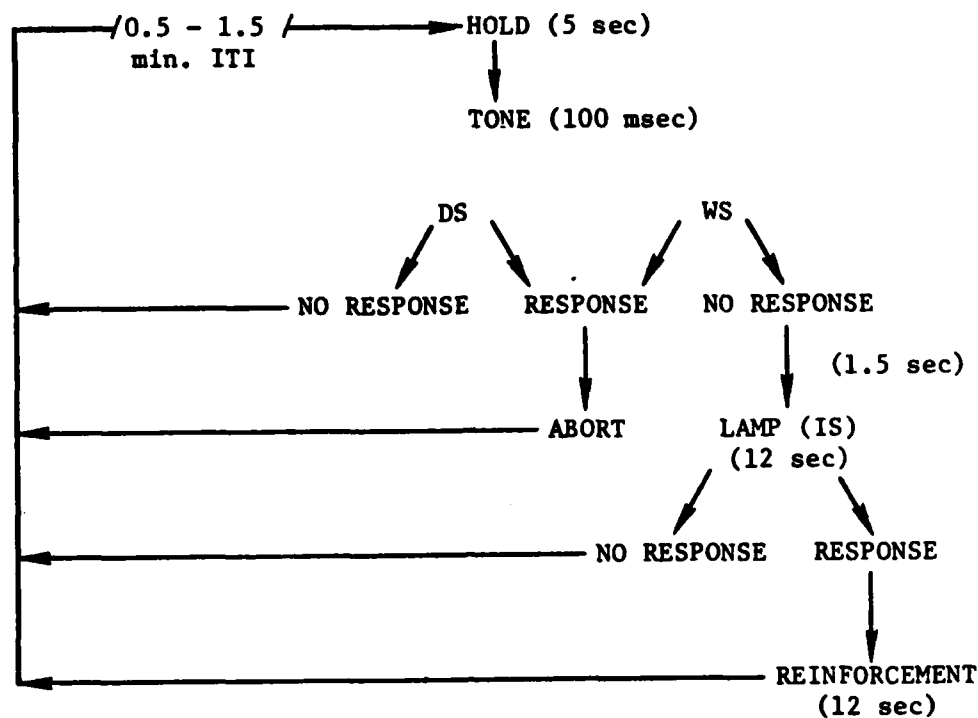


FIGURE 1 LOGIC OF THE CUED REACTION TIME TASK

aborted if a premature response occurs, and the IS terminates if a response is not made quickly enough. The logical configuration of this system is presented in Appendix A.

Associated with the Iconix unit were three counters that measured reaction time and counted the total number of bar presses and reinforcements. Liquid was delivered through a solenoid, which gated a gravity or pressurized flow system so that 1 ml of juice was delivered with each bar press.

Several limitations of this preliminary system mandated an upgrading of the facility. These limitations included the following:

- A hard-wired system is inherently inflexible; therefore, new wiring configurations have to be done to implement even trivial experimental changes.

- The logical control system was only indirectly tied to the electrophysiologic data-acquisition system; so automatic tagging of data, automatic sorting, and selective averaging could not be accomplished.
- Behavioral data had to be tabulated manually.
- Tone frequencies had to be changed manually, requiring a manual system for keeping track of WS and DS trials.
- Summary statistics of behavioral data could not be obtained immediately after a test session.
- The gravity and pressure flow systems were cumbersome and messy.
- Obsolete Linc-8 computers were inadequate for modern requirements of data analysis.

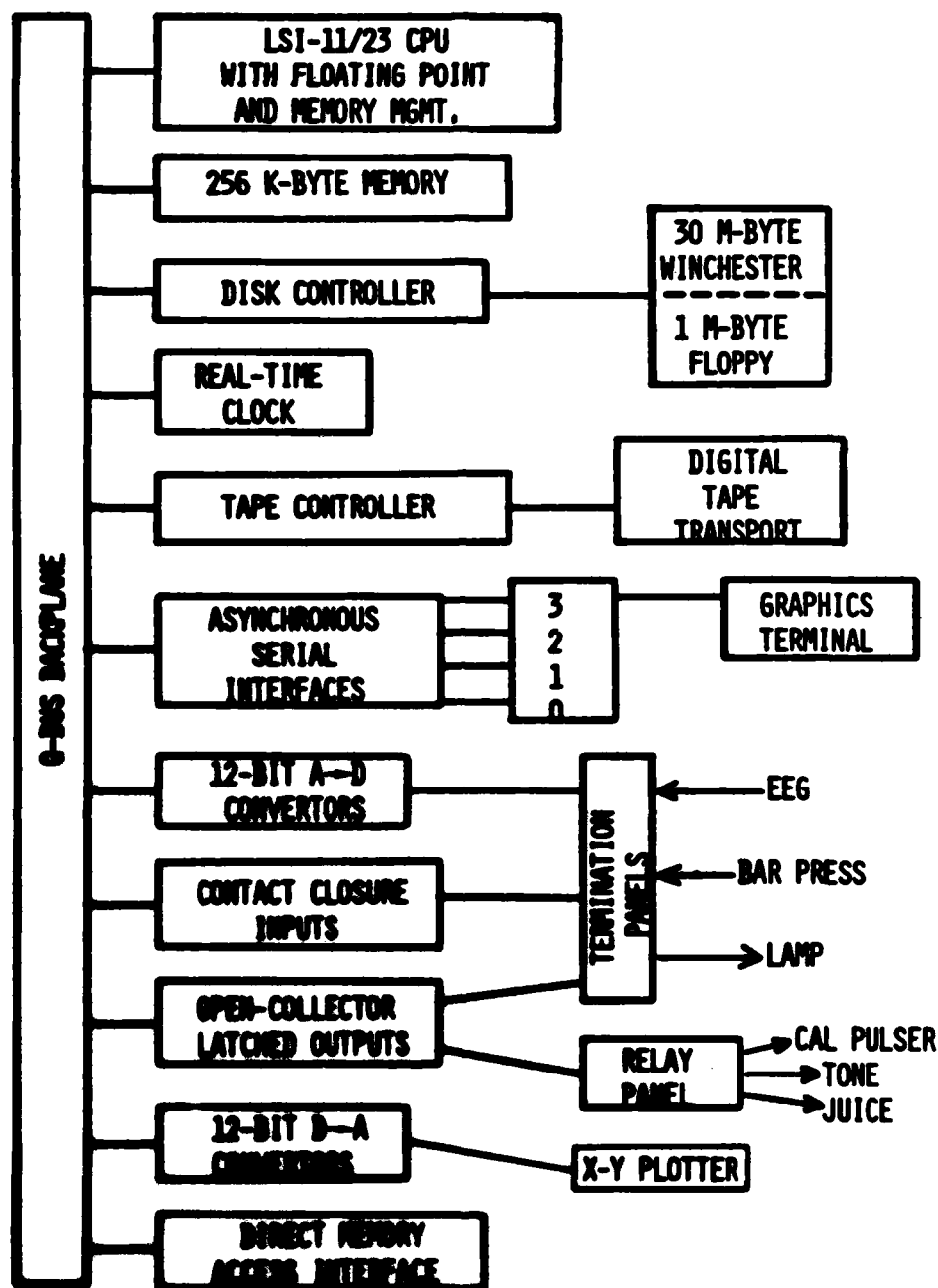
Consequently, SRI purchased the LSI-11/23 computer system described below for use on this project.

### 3. LSI-11/23 Computer and Related Components

#### a. Hardware Configuration

The configuration of this system is schematized in Figure 2. It consists of an LSI-11/23 processor with extended memory (256 KB), clock board, analog-to-digital converter and associated direct memory access board, a digital-to-analog converter, contact closure detector, latched open-collector board for operating external devices, a 30-MB Winchester disk with associated 1-MB floppy, a 9-track digital tape recorder, and VT640 graphics terminal. Associated devices include solid-state tone generators under computer control, a circuit interface between the computer and liquid delivery system (a Valcor 5P94R-7 metering pump), a gain and DC-offset control panel, indicator panel, H-P model 7034A X-Y plotter, and TTY Model 43 printer.

The Winchester disk is used to store programs and, temporarily, single-trial data during testing. At the end of the test session, the single-trial data are transferred to digital tape. The floppy discs are used to store waveform averages and summary statistics of behavioral data for the session. The summary statistics are printed on the TTY-43 printer at the end of the session, and waveform averages are plotted on the HP plotter. Listings for the programs developed for this system are provided in Appendix B.



**FIGURE 2 SCHEMATIC OF COMPUTER SYSTEM**

The central processing unit is a digital equipment corporation Model LSI-11/23 with extended memory and floating point enhancement. It's Q-Bus backplane holds an array of special-purpose hardware boards to operate peripheral devices.

b. Software Development

CNV paradigm. A program was written to implement the logical paradigm shown in Figure 1. The program was designed to provide all record-keeping and summary statistics of behavioral data and to execute the paradigm while obtaining the ERP data. Thus, the parameters of stimulation and recording, sampling rates, artifact-reject criteria, experimenter remarks, ratios of WS and DS trials, date, experimental conditions, and so on are stored with each set of waveforms so that manual notations and paper records are minimal.

Program "CNV" implements a cued RT paradigm by controlling the sequence of stimulus events, acquiring data from each of up to 8 electrode input channels, monitoring responses by the subject (S) and input from the experimenter (E), and labeling and storing all results on Winchester and floppy disks.

E begins each session by calling up "CNV" and responding to its queries regarding:

- Name of the subject
- Date
- Experimental condition and day within the condition
- Channels that are to be sampled.

Also, E has a chance to modify the values of various trial parameters:

- Probability that each trial will have a WS rather than a DS tone
- Longest RT following light (IS) onset that will still be rewarded
- Smallest voltage that will be considered an "artifact"
- Channel to be automatically displayed at the end of each trial
- Amplitude of the calibration pulse for each channel
- Interval between tone onset and light onset
- Duration of the IS light
- Speed mode--i.e., whether optional ITI activities are conducted.

Each trial is begun by E's pressing a button when the computer, E, and S are all ready. Seated at the computer terminal, E can watch S on closed-circuit TV, supervise the flow of the computer program, and access the polygraph or other equipment, as needed. Trial-by-trial results are reported on the terminal screen, including the number of bar presses by S, reaction time, duration of ITI, and number of ITI responses.

The program logic considers each trial to consist of seven consecutive "phases," demarcated by experimental events: onset and offset of a tone, onset of the IS light, termination of sampling, etc. To time both the sampling (one set of A-D conversions every 10 msec; 400 consecutive sets per trial) and the stimulus events, a clock is set to overflow once per millisecond. Every 10th overflow, sampling is executed and the obtained values are stored in a buffer. Between samples, a contact closure register is polled to see if any bar presses have occurred. If S presses the response bar while the IS is on, then the metering pump is operated to provide a juice reward to S. Bar presses at earlier times during a trial are tabulated according to the trial phase in which they occur; also, they cancel the IS and the opportunity to earn juice. If there is a contact closure--on a separate circuit--due to E rather than S, an ongoing trial is aborted.

After checking for contact closures, "CNV" evaluates whether it is time to begin the next phase of the trial, as follows:

<u>Phase</u>	<u>Time (msec)</u>	<u>Event</u>
1	0	Begin sampling
2	400	Begin trigger for CAL pulse
3	450	End trigger for CAL pulse
4	1000	Turn on tone
5	1100	Turn off tone
6	2500	Turn on IS (if WS trial)
7	4000	Stop sampling

On WS trials, unless S has bar-pressed prematurely, the IS appears during Phase 6 and the reaction time to bar press is recorded. (On DS

trials, although no IS is presented, responses are nevertheless recorded for all seven phases of the trial.) If a bar press happens within a 3000-msec, "limited-hold" interval after IS onset, the IS remains on for a total of 12 sec and each subsequent bar press is rewarded. If no bar press is made in time, the IS ends and the ITI begins. At the end of each trial, E has a chance to insert a comment that will be stored with the data of the previous trial.

To detect bar presses during the ITI, the contact-closure register is set so that all bar presses are noted (including when they occur), meanwhile enabling the system to perform other tasks. Data from the foregoing trial are screened for artifacts, and their maxima, minima, and DC levels are computed. All this information, along with E's comments, settings of all parameters, and the waveform data, is written--as a single record--into the session's file on disk.

Next, the waveform data are added into an averaging buffer; separate buffers are used for data from WS vs. DS trials. However, if there were too many artifacts or if the trial was aborted (due to premature bar press, too slow RT, or E's intervention), then the data are not added to these buffers.

Unless E has opted for "high-speed" mode (short ITIs), waveform data from one of the channels is displayed on the graphics terminal. E can opt to review single-trial data from any channel or cumulative averages for WS or DS on any channel. When E is finished displaying data, he then selects the next task, from a menu:

Hit RETURN for next trial, or select item from menu:

??---Explain menu  
QT---Quit session; delete data  
EX---Exit session; save data  
DS---Display single-trial data  
DA---Display average data  
CP---Change parameters  
PL---Plot waveform

Unless the session is over, E ordinarily waits out a variable ITI and then proceeds to the next trial. Occasionally, to review/modify the trial parameters, E selects "CP" and then, interactively, changes any or all (or none) of them. If a hard-copy of a waveform is needed, "PL" initiates that interactive sequence.

When the sample sizes for WS and DS trials are adequate (e.g.,  $n > 15$ ), E selects "EX" from the menu. This causes the session's single-trial data and information to be saved as a file on the Winchester disk. Later, E will transfer this file to magtape. Next, a summary is prepared of the statistics and parameters for the session, including:

- Total number of trials
- Sample sizes for WS and DS trials
- Number of trials aborted for each possible reason
- Mean RT
- RT standard deviation
- Mean number of reinforcements
- Total number of bar presses
- Number of bar presses during ITIs
- Number of artifact-rejected trials
- Mean number of artifact-rejected trials
- Mean number of deviant data per artifact-rejected waveform per channel

The WS and DS averages, along with the session's parameters and statistics, are stored as a file on floppy disk. Finally, a session summary and a listing of E's comments are output on a line printer.

P300 paradigm. A program was written to carry out preliminary studies of the P300 evoked by tones of different frequencies in a "passive" paradigm--i.e., no responses are required of the monkeys. However, when the program was used with human Ss, the Ss were asked to count the rare tones because we were primarily interested in verifying the adequacy of the program.

Program "P300" implements a P300 paradigm in which 100-msec tone pips are presented at a fixed rate (e.g., once per second). Differently

pitched tones are included in each series; "common" tones occur more often than "rare" tones. No overt response by S is required.

Four channels of the A-D converter are used to sample potentials from various electrode sites. Each "trial" consists of an epoch of 1000 msec during which 400 samples per channel (4 channels) are acquired, one set of samples every 2.5 msec.

To begin a P300 session, E calls up "P300," enters the name of S, and inputs a name for the file to which data will be written. Then, E has a chance to interactively modify any of the following parameters:

- Percentage of common tones
- Minimum value that will be considered an "artifact"
- Amplitude of each channel's calibration pulse
- Duration of recording epoch
- Interepoch interval
- Speed mode--i.e., whether optional activities occur between epochs
- Number of the upcoming tone series.

At the beginning of each tone series, E decides whether the more highly pitched of two tones will be common or rare. E then specifies how many trials are desired. The sample size for the rare tone is computed so that it can be used as a criterion for completing a tone series. Subsequently, if the waveforms for some rare tone epochs are artifact-rejected, the series will be continued until the sample size for the rare tone is adequate. When E, S, and the computer are all ready, E begins the tone series by pressing a button.

On each trial, as sampling is begun, a relay is used to trigger a delayed calibration pulse. About 200 msec later, one of two relays is operated--to deliver a common or rare tone, depending on the output of a random number generator and on the percent of common tones previously requested by E. Sampling continues for about 625 msec after the offset of the tone.



Between samples (every 2.5 msec), the values produced by each set of A-D conversions are stored in a buffer. Then the time is checked to see if the next event in the trial is due. After the 400th set of samples is in, all the data are reviewed for artifacts while their maxima, minima, and DC levels are being computed. Unless too many artifacts occurred, the data are added to another buffer, for averaging; separate buffers are used for the common and rare tone data. If more rare tone samples are needed, the next trial begins as soon as the interepoch interval has elapsed. Between trials, E is informed of the tone type for the next trial, the current sample sizes, and the DC levels.

Following the completion of each tone series, E has a variety of options: to add more trials and extend the tone series; to write the data--along with parameter information--as a record in the session's file; to begin the next tone series; to list the current parameter values; to display either the common or rare tone averages (or the most recent single-trial data) for any of the four channels; to plot any of these waveforms on an X-Y recorder; to get a status report on the just-completed tone series; or to end the session.

As each set of data is written to disk, a summary of the experimental conditions is printed out. It includes items such as:

- Date and name of subject
- Number of preceding tone series
- Name of data file on floppy disk
- Amplitudes of calibration pulses on each channel
- Common vs. rare tone sample sizes
- Total number of trials
- Numbers of common vs. rare tones presented
- Smallest voltage value to be deemed an "artifact"
- Number of samples artifact-rejected
- Time between samples
- Time between epochs.

Data retrieval program. Data analysis software is being developed that will enable us to reopen any data file written during a CNV, P300, or other class of experiment, recall what conditions were employed, review the data obtained, and sort or analyze the waveforms as needed. Currently, we are able to display waveforms off-line on our graphics terminal and "score" the voltage values at any point along the waveform. With a graphics printer slaved to the terminal, we will soon be able to obtain report-quality hard copies of anything (text, graphs, bar charts, or waveforms) that can be displayed on the terminal. In the meantime, we are operating an analog X-Y plotter with our D-A converter to plot waveforms recorded during pilot experiments. A line printer is used to print out textual information.

### C. Verification of System Performance

Electrodes somewhat different from those used previously for recording subcortical slow potentials (SPs) from monkeys are now being employed (see Section E, Surgical Preparations; briefly, the pipettes were sharpened so that multiple units as well as SPs could be obtained), and the low-frequency filters on the EEG amplifiers (Grass Model 7P511) were modified to 0.01 Hz to allow measurement of SPs. Therefore, recordings were made from the lateral geniculate of one acutely prepared cat to check these items. Because of the fairly high electrode impedance, a major purpose of the check was to ensure that recordings without excessive noise could be obtained.

The cat was anesthetized with sodium pentobarbital (42 mg/kg, i.p.) and placed in a stereotaxic instrument. An incision was made in the scalp, and fascia was cleared from the skull. A small hole was drilled in the skull at coordinates appropriate for approaching the lateral geniculate nucleus (LGN), and a saline-filled pipette with a 100- $\mu$  tip was lowered into the LGN while flashes from a Grass PS-2 photostimulator were being presented.

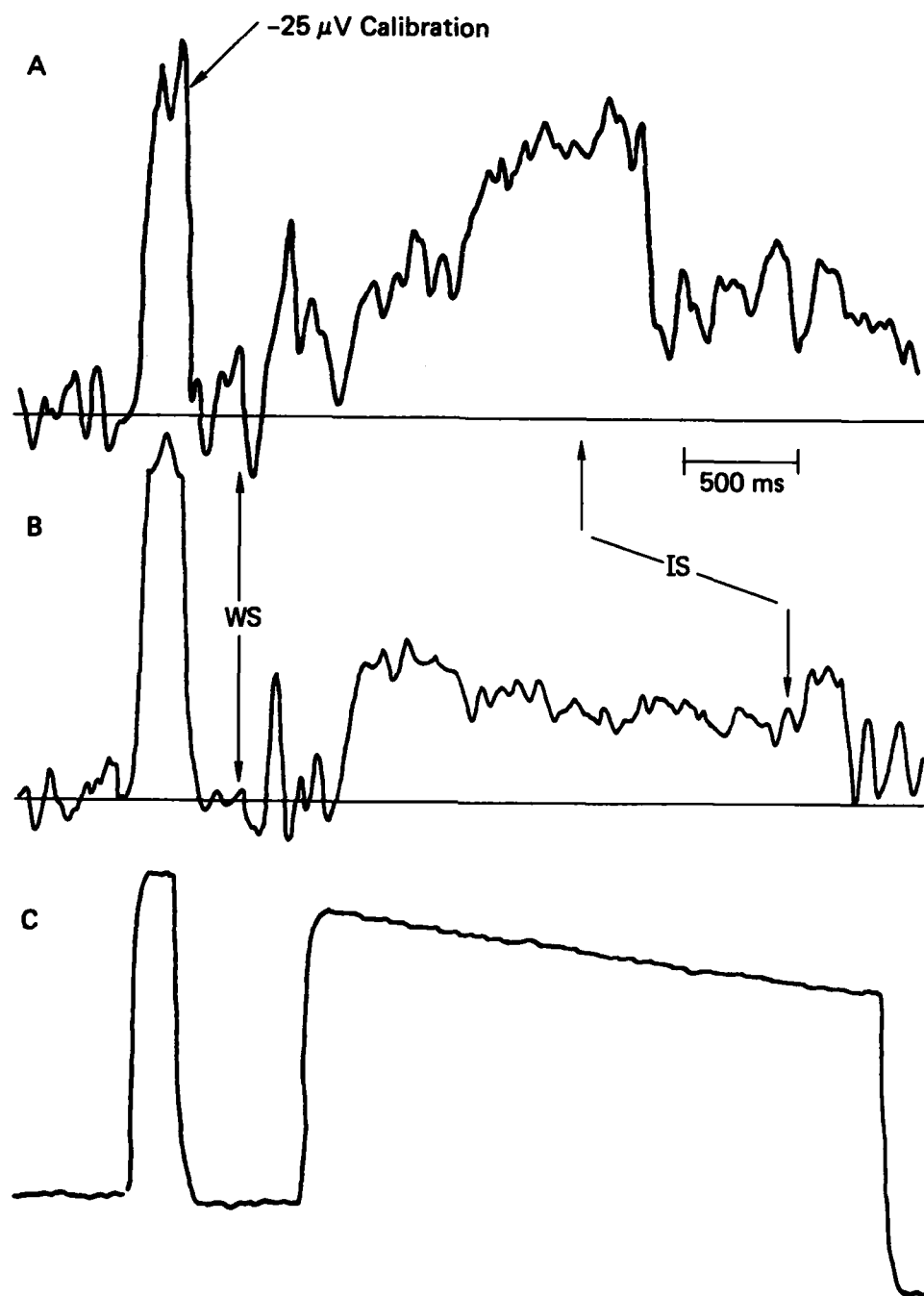
We could not conveniently average the ERPs at this time, but multiple-unit activity comparable to that observed in earlier studies

(Rebert, 1973) was obtained and slow potential ERPs could be observed sufficiently frequently in the raw records to confirm the adequacy of the amplifier low-frequency modification. Thus, both slow and massed-unit responses were obtained through the same electrode and amplifier with adequate resolution, except that the 60-cycle filter had to be used for ERP recording.

Several human Ss were tested in both the CNV and P300 paradigms to evaluate the computer program performance as well as the amplifiers. Beckman Ag-AgCl discs were placed on the mastoids for reference and above and below the left eye for recording the vertical electrooculogram (EOG). A stretchable cap containing a conductive gel-filled tube attached to another Ag-AgCl disc was used to obtain vertex records. The forehead was grounded via another Ag-AgCl disc. The warning and discriminative tones, the imperative stimulus (light), and the response manipulandum to be used with the monkeys were also employed to test the human Ss. The same tones were used in the P300 paradigm, and Ss were instructed to count the rare tone.

Figure 3 shows CNVs from two Ss; the first was tested with an interstimulus interval (ISI) of 1500 msec, and the second with an ISI of 2500 msec. Recording was with one of the Grass 7P511 amplifiers modified to have a low-frequency cutoff of 0.01 Hz. These are very typical CNVs. The CNV in the long ISI exhibits some decay consistent with the amplifier time-constant shown in the lower part of the figure. We determined that this time-constant was consistent across the eight channels to be used with the monkeys.

Examples from one S of P300 responses associated with several proportions of rare tones are shown in Figure 4. The inset shows a typical decline in N200-P300 amplitude as the proportion of rare tones increases.



**FIGURE 3** CNV AND SQUARE WAVE RECORDINGS WITH 0.01 to 100 Hz BANDPASS

A. CNV from one S with 1500 ms interstimulus interval. B. CNV from another S with 2500 ms interstimulus interval. C. Square wave showing time-constant of the recording system.

% RARE TONE

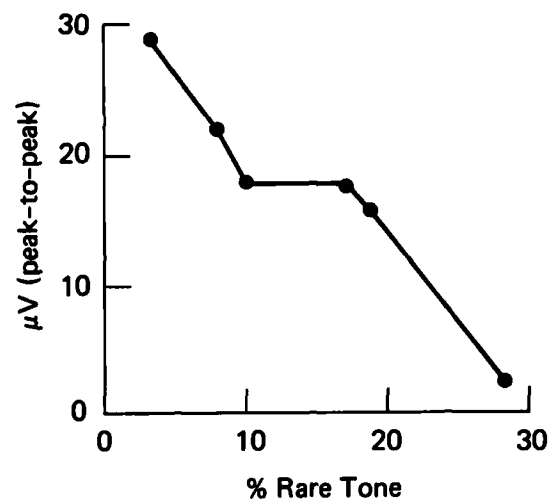
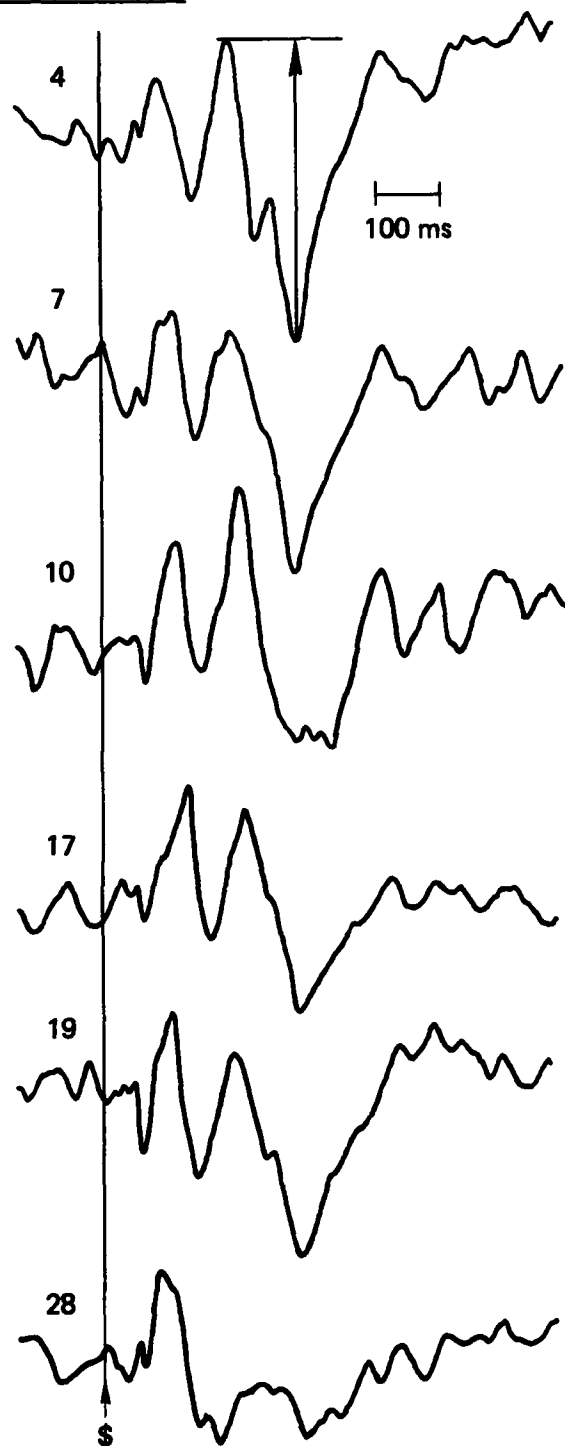


FIGURE 4 EXAMPLES FROM ONE S OF VERTEX POTENTIALS EVOKED BY RARE TONES IN AN "ODDBALL" TASK, AND THE PEAK-TO-PEAK P300 AMPLITUDE AS A FUNCTION OF THE PERCENTAGE OCCURRENCE OF RARE TONES

#### D. Preliminary Activities with Monkeys

##### 1. Quarantine, Housing, and Care

Six adult male cynomolgus monkeys were received on 10 June 1982. In accordance with legal requirements, the monkeys were quarantined for one month. During this time they were weighed periodically and tested for communicable diseases and general health. One monkey exhibited anemia, lack of appetite and diarrhea for about a week. Mean individual weights of the six monkeys during quarantine ranged from 4.3 to 6.1 kg. The individual weights were very stable during this time. Housing was in standard monkey cages with automatic watering. Chow was given twice per day.

##### 2. Preliminary Training

The monkeys were released from quarantine on 20 July 1982. Chair training for three monkeys was started on 30 July. Two monkeys could not be handled because of their aggressiveness and large canines. The canines were cut on 6 August. During this time we were able to train only four monkeys because considerable time was devoted to general handling, taming, and inhibiting the fierce aggressiveness of one monkey (Samurai). During the first week of August, juice, raisins, grapes, and bananas were given to the monkeys while they were chaired. Reactions were highly mixed; M22 (E.T.) ate or drank anything proffered, but M24 (Mickey) would partake of nothing. Although quite aggressive and noisy while being handled, all the monkeys were quiet when chaired, exhibiting very little struggling or turning.

##### 3. Operant Conditioning

During training sessions a monkey was placed in a primate chair with some special features. A drinking tube was mounted so that it was readily accessible to the monkey, and an inverted, metal U-bolt covered with tygon tubing arched over the monkey's snout and prevented turning of the head. A Plexiglas partition to the left of midline of the belly plate kept the arms separated and encouraged the pressing of the response bar with the right hand. The bar--a 5-cm wide piece of

stainless steel--protruded 5 cm from the Plexiglas plate on the front of the chair, 27 cm from the monkey's stomach. A ball mounted on the midline of the belly plate 12 cm from the monkey's stomach provided a place for the monkey to put its right hand prior to a trial in order to stabilize reaction times. A light box (4.5 x 4.5 cm square) with a white plastic front was mounted on the chair 27 cm away from the monkey's brow and 10 cm above eye level. Intensity of the light, which constituted the IS, was 400 ft-lamberts. Two video cameras and monitors were used to obtain (1) a wide-angle view of the monkey and (2) a close-up of the ball so that trial initiation by E could be made contingent on a proper placement of the monkey's hand.

Training proceeded in three phases: (1) bar-pressing for reward, (2) light-contingency, and (3) pretrial placement of the hand. During Phase 1 successive approximations of the bar-press response were reinforced until the bar was consistently pressed. In Phase 2 the delivery of reward was made contingent on presence of the IS. This involved extinguishing responses during dark periods and training the monkeys to tolerate widely varying interstimulus intervals so that they would be responding to the IS rather than in a temporal pattern. Phase 3 was initiated during early stages of Phase 2--when the monkeys indicated some awareness of the meaning of the light. This phase involved using the IS as a secondary reinforcer to reward placing the right hand on the ball. Later in this phase, RTs were recorded and a limited-hold contingency was included so that slow RTs resulted in extinguishing of the IS and unavailability of reward. This procedure is included so that, if necessary, the RT can be pushed to the point where use of the warning signal (WS) is necessary in order to have an RT sufficiently short to obtain reward (there is nothing else in this paradigm that requires the monkeys to pay attention to, or utilize, the WS, although in previous work they did so).

Bar-press training was initiated with E.T. on 9 August and he almost learned the response on that day--he did learn that a reaching movement was required, and the next day he began to bar-press. By 16

August, M23 (Conan) had also learned to bar-press and the two others (Samurai and Smacker) were making arm movements to precipitate a reward. Large individual differences in motivation were evident at this time. E.T. would press immediately and rapidly upon availability of the bar, registering, for example, 849 responses in 15 min. However, neither Samurai nor Smacker was very motivated. Conan exhibited intermediate behavior, pressing about 450 times in 15 min. By 26 August, Samurai and Smacker were partially on the bar, pressing occasionally but they subsequently regressed and would not perform. These monkeys were then deprived of all water and were required to satisfy their liquid needs in the experimental environment. Smacker responded to this satisfactorily and began to bar-press consistently at 200-300 responses in 15 min. Samurai, however, was extremely erratic and developed a peculiar pattern of not responding unless an E was inside the chamber with him. We concluded that this monkey might not be satisfactory in the long run. E.T. was started on light-contingency training on 27 August and progressed satisfactorily.

Monkeys 24 and 25 (Mickey and Grey) were first chaired on 13 September, having been liquid-deprived for 48 hr. Although Mickey was very calm in the chair, he was not the least interested in drinking. In contrast, Grey was well motivated and learned to move his arm to receive reinforcement. He again exhibited diarrhea and was not studied until 28 September, when he learned to bar-press moderately well.

We discovered that the monkeys would lick and pick small drops of water off their cages whenever the cages were rinsed. This made no difference to either E.T. or Grey but was sufficient to make Mickey totally uninterested in the experimental situation. We then ensured that the cages were dried before the monkeys were replaced.

Training continued through 8 December, progressing through the several phases until all the monkeys were placing the hand on the ball and exhibiting relatively short RTs. They were then tested only on Mondays for three weeks, with liquid deprivation only over the weekend, to determine whether they would retain their performance after such



layoffs, which would occur during the later test sessions following electrode implantation. Performance was maintained. On the last day of testing, mean RTs over trials ranged, among the monkeys, from a low of 894 msec (SD = 164) to a high of 1812 msec (SD = 950). The rate of bar-pressing ranged from 1.0/sec (SD = 0.1) to 1.7/sec (SD = 0.2). For the five monkeys consistently responding (all except Samurai), mean RT was 1389 msec (SD = 342, SEM = 153) and the mean rate of pressing was 1.4/sec (SD = 0.3, SEM = 0.1).

#### E. Surgical Preparations

##### 1. Placements

Preparations for surgery were initiated during the last week of December 1982. They involved calculation of electrode placements and construction of electrodes. Because no stereotaxic atlas was available for the size of cynomolgus monkey (Macaca fascicularis) being used, estimates of placements were determined from atlases for M. fuscata (7.4 kg) and small M. fascicularis (3.5 kg) and from sections of M. arctoides from a previous study. These last sections were critical to a determination of placement depths because other atlases do not show the cortical surface, and the dura must be used for M. fascicularis because of extreme interanimal variability of brain placement with respect to the stereotaxic landmarks--auditory meatus and inferior orbit (Dubach and Bowden, unpublished manuscript).

Because we were unable to study the number of monkeys originally proposed, the strategy for studying different placements was altered. We decided to put electrodes bilaterally over premotor (arcuate region), motor, and parietal cortices, in the CA1 region of the hippocampus, and in the caudate nucleus, substantia nigra, midbrain reticular formation, and n. ventralis anterior of the thalamus. Two references were used in case one became dysfunctional; these were placed in anterior and posterior white matter. An electrode was also implanted in the superior bony orbit to record the EOG.

## 2. Electrodes

Electrodes capable of simultaneously recording DC potentials, transient ERPs, and massed-unit activities were constructed of glass pipettes and cells housing commercially available sintered Ag-AgCl pellets. These were like the electrodes described by Rebert and Irwin (1973). The tips were sharpened to 100  $\mu$  by pulling 0.7-mm O.D. pipettes to very small tips and then, under a microscope, chipping away the tip to the 100- $\mu$  size. This was facilitated by first etching a 100- $\mu$  diameter platinum wire to a gradual taper and inserting it into the pipette. The wire provided a means of determining when the tip diameter was the right size. A short piece of shrink tubing was applied to the upper shank of the pipette to prevent its slippage in the stereotaxic electrode carrier during surgery. The pipettes were then filled and sterilized by boiling in normal saline.

Electrode cells, which are placed over the top of the pipette during surgery, were constructed of 1-cm lengths of 4-mm O.D. glass tubing. One end of each tube was fire-polished to an inside diameter of 1 mm (just large enough to slip over the pipette). These tubes were half-filled with Agar-saline followed by normal saline when the agar had set. A Ag-AgCl pellet was then placed in the tube, resting on the agar, and the upper opening was closed with dental acrylic. These cells were stored in a freezer until used.

## 3. Surgery

Monkeys were deprived of food the night before surgery. Ketamine hydrochloride, a fast-acting nonbarbiturate anesthetic, was administered intramuscularly (i.m.) at a dose of 14 mg/kg, which was sufficient to heavily tranquilize the monkey for preparatory activities. A dose of atropine sulfate (a 0.2-ml solution of 120 g/ml) was administered i.m., the head and calves were shaved and cleansed, and then the monkey was placed in the stereotactic apparatus. A catheter was placed in the right small-saphenous vein, through which sterile saline was dripped at a rate of 16 drops/min--sufficient to keep the catheter clear and provide some hydration during the 4-5 hr of surgery. The

electrocardiogram (EKG), and the electromyogram (EMG) of the right triceps, were recorded. The EMG was used to monitor state of alertness, and 0.2 to 0.3 ml of sodium pentobarbital (65 mg/ml) was administered through the catheter as required. Typically, an infusion was required each half-hour. Rectal temperature was also monitored; it was very stable during surgery and ranged from 32.5 to 34.5°C in the six monkeys.

Muscle and fascia were cleared from the skull and the small blood vessels were cauterized. Then, stereotaxic coordinates for the anterior-posterior and lateral planes were marked on the skull. Four 6-32 stainless-steel bolts were threaded into burr holes, and bone primer and a layer of dental acrylic were applied. The EOG electrode was inserted into a burr-hole in the frontal bone, then the remaining holes were drilled. After the stereotaxic reading for the dura was obtained the electrodes were lowered into the brain. The electrode cells were then attached and wired to a 14-pin plug. The assembly was encased in acrylic, the wounds were sutured, and 0.5 ml of penicillin-G in dihydrostreptomycin sulfate was administered i.m. Furacin (nitrofurazone) antibiotic salve was applied around the headplug.

Antibiotic was given for 2 to 5 days postsurgically. Each day, the head was washed, and the wound was infiltrated with Liquimast (oxytetracycline HCl).

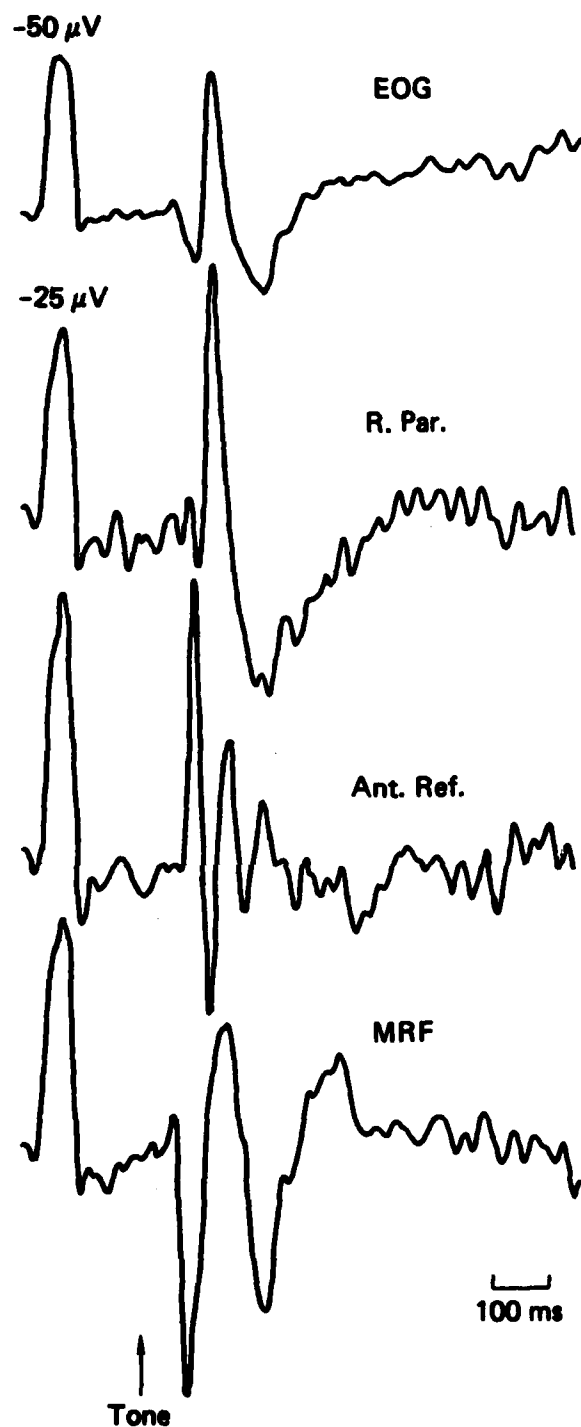
#### F. Electrode Tests in Monkeys

Several weeks following surgery, the electrodes of each monkey were tested one at a time with a DC amplifier to check offset, drift, and noise. The monkey was restrained in the chair but the manipulandum was not present and no stimuli were given during the electrode checks.

DC offsets were usually less than 10 mV and were mixed with respect to polarity. Four electrodes in E.T. were 13.0, 13.2, 14.0, and 14.8 mV, and two in Smacker were 21.3 and 11.3 mV. Drift was generally negligible in terms of the 4-sec sampling epoch for the CNV program and expected amplitudes of slow potentials (drift less than 20  $\mu$ V/sec, with expected amplitudes of several hundred  $\mu$ V). Drift will be attenuated by

the capacity-coupling to be used. Occasionally, drift was much as 50  $\mu\text{V}/\text{sec}$  at the beginning of the test, but this subsided with continued recording--a common characteristic of Ag-AgCl electrodes. Most of the subcortical electrodes picked up 60-cycle line noise, requiring the use of the 60-cycle filter on the amplifiers. In two monkeys two and four of the subcortical electrodes were very noisy and possibly open for unknown reasons (perhaps bubbles in the pipette). We are in the process of constructing cable from microdot wire and rearranging the laboratory cabling to improve the recording environment for these high impedance electrodes. If necessary, preamplifiers with very-high-input impedance will be employed, or more drastic filtering of the EEG (e.g., with a low-pass setting of 40 Hz) will be used.

After the preliminary electrode tests had been completed, the WS and DS tones were presented to observe tone-evoked potentials. Examples from one of the monkeys are shown in Figure 5. For these examples the EOG, a cortical placement (right parietal), subcortical nucleus (midbrain reticular formation), and subcortical white matter (anterior reference) were used. The cortical placement showed a response very similar to that recorded in the EOG channel, and there were similarities to the EOG in the other placements as well, although the number, latencies, and polarities of components were different than would be expected if they were just EOG artifacts. These transient potentials are obviously recorded from intracerebral tracts (anterior reference) as well as from the nuclei. That the EPs recorded were probably not EOG artifacts is suggested by Figure 6, which shows the raw EEG tracings from these placements in the absence of tone pips. Whereas the EOG channel exhibits blinks of about 250  $\mu\text{V}$  and eye movements, these events are not clearly present in the EEG records, although there are some later-occurring shifts that might be neural events time-locked to the blinks (e.g., the slow shift in parietal cortex). The slow eye movements have little or no effect on the recordings. To more thoroughly examine the effects of blinks and eye movements on the EEG records, we will use a continuous sampling routine, which is terminated



**FIGURE 5** TONE EVOKED POTENTIALS FROM SUPRAORBITAL RIDGE (EOG), RIGHT PARIETAL CORTEX (R. PAR.), ANTERIOR WHITE MATTER REFERENCE (ANT. REF.), AND MIDBRAIN RETICULAR FORMATION (MRF), WITH RESPECT TO THE POSTERIOR WHITE MATTER REFERENCE

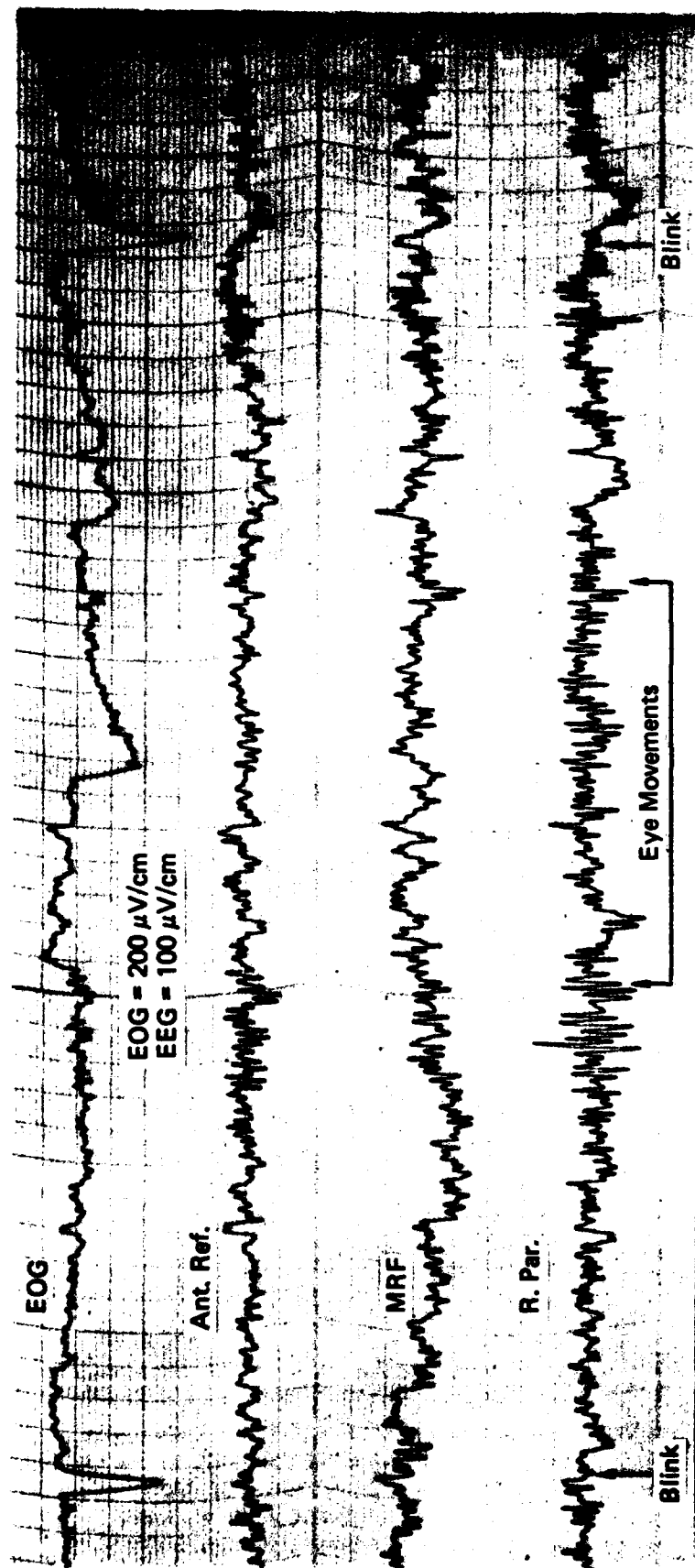


FIGURE 6 EEG RECORDS FROM ONE MONKEY SHOWING MINIMAL EFFECTS OF BLINKS AND EYE MOVEMENTS ON CORTICAL AND SUBCORTICAL RECORDINGS

EOG = electrooculogram; Ant. Ref. = reference in anterior white matter; MRF = midbrain reticular formation; R. Par. = right parietal cortex. All recordings, including the EOG, were made with respect to the reference electrode in posterior white matter.

on detection of a blink so that averages time-locked to the blinks can be obtained.

#### G. P300 in Monkeys in Response to Visual Cues (Stanford University)

##### 1. Background

Of the several transient potentials of the brain associated with the perception and processing of environmental events, the P300 "component" (a complex of late positive waves) is the most clearly associated with cognitive events. This "component" seems to reflect only the general informational properties of stimuli and is uninfluenced by informationally irrelevant physical characteristics or by specific behavioral response requirements (Pritchard, 1981). These characteristics have made the P300 complex particularly relevant to the evaluation of workload, to the general field of biocybernetics (e.g., Donchin et al., 1982), and to clinical research on cognitive disorders (Hillyard and Kutas, 1983). Consequently, an interest in obtaining a more complete understanding of the intracerebral sources of the P300 complex has developed (Galambos and Hillyard, 1981). However, to date the results of research provide only hints about the electrogenesis of the P300. For example, Halgren et al. (1980) recorded P300-like activity in amygdala and hippocampal gyrus, Wood et al. (1980) noted the lack of polarity reversals of P300-like events in depth recordings above the hippocampus, and Okada et al. (1983) concluded from magnetic field recordings that the P300 is generated in anterior hippocampus. On the other hand, Johnson (NIMH, personal communication) has recorded the P300 complex from human patients with hippocampal lesions. But there is little concrete evidence localizing the sources of the P300 phenomenon and, clearly, invasive studies in animals are necessary to systematically study this point.

##### 2. Facilities and Equipment

A behavioral testing system used extensively in a variety of neuro-behavioral studies of monkeys at Stanford University (Kimble et al., 1965; Bagshaw et al., 1965; Grandstaff and Pribram, 1971; Pribram

et al., 1980) is being employed in this investigation. The system hardware consists of PDP-11/34 and Apple computers implementing a "discrimination apparatus for discrete trial analysis - sixth version" (DADTA-VI--Cutcomb et al., 1981). Nine translucent panels in front of a color television set, controlled by the Apple II microprocessor, serve as both stimulus display and response panels. The PDP-11/34 controls the Apple II and records panel press position, reaction time, error scores, and electrical brain activity during task performance. During testing, a monkey is restrained in a primate chair housed in a modified refrigerator. Correct responses result in the delivery of liquid reinforcement.

### 3. Test Paradigms

Two "oddball" procedures--behaviorally nondiscriminative and discriminative--are being implemented. Although stimulus relevance is an important parameter in P300 genesis, the attribution of significance to a stimulus by requiring an overt response leads to confounding of motor and cognitive processes. Thus, in the first task being studied, no differential responses to rare and common stimuli are being required. After the monkey presses a central panel to initiate a trial, one of the remaining eight panels (randomly selected) lights up and the monkey must press the lighted panel. The response panel lights red a designated percentage of the time and lights green on other trials. Although this paradigm involves behavioral responses, it is equivalent to passive tasks shown by others to produce P300-like events in monkeys (H. J. Neville; S. A. Hillyard; personal communications) in that the two stimuli provide no differential instructions to the animal with respect to performance. Nevertheless, we expect the color novelty to elicit a P300-like potential. In a second task the monkeys will respond to only one color and the proportion of trials on which that color occurs will be varied. We expect P300-like potentials to occur to either the red or green stimulus when it is the rare event, but the potential should be larger in response to the behaviorally relevant stimulus when it occurs only occasionally than to the nonresponse stimulus when it is rare



(Hillyard and Kutas, 1983). In addition, the cortical and subcortical distributions of the "P300" complex will probably vary as a function of those manipulations.

#### 4. Electrodes and Placements

Electrodes like those described by Barna et al. (1981) are being constructed. The electrode consists of 30-gauge stainless-steel tubing, tapered at the tip and housing a maximum of 16 electrode wires. The desired number of wires are tied into a bundle and threaded through the tubing and fixed in place with epoxy. When this has partially set, the wires are stripped from the bundle, bent at right angles to the shaft, and cut flush with the outer surface of the tubing. The surface is then smoothed to form a pointed shaft. This array provides the capability for searching out dipoles indicative of generator sources and recording massed-unit activity, which is also helpful in source localization (Vaughan, 1981). These types of electrodes, with varying numbers of leads, will be placed in the three major nuclei of the amygdala, in the caudate nucleus, and in the anterior hippocampus and dentate gyrus.

#### 5. Preliminary Behavioral Results

Two female M. fascicularis monkeys have been trained on initial parts of the "passive" P300 task described above. Training proceeded in four phases: (1) chair taming, (2) trial initiation, (3) sequential start and panel-pressing, and (4) color cue contingency. During Phase 1 the monkeys received apples and Tang® while in the chair. This training occurred on most weekdays for about a month. Behavioral training in Phase 2 involved teaching the monkeys to press the central panel of the 3 x 3 matrix when a small white circle appeared. This stimulus-response sequence ensures that the monkey is attending to the display panel. During Phase 3, presses on panels other than the central one, following the central press, were rewarded. This training also involved making only a subset of the panels relevant to prevent position habits. Finally, in Phase 4 the animals were trained to make the second press on the panel behind which a green square was presented and response latencies were required to be less than 3 sec. Variation of latencies

between the center press and lighting of the second panel was also introduced to reduce any tendencies to "time" the interval and produce unusually low reaction times. The number of trials per session was extended beyond 100 as the monkeys became used to obtaining most of their daily liquid in the testing situation.

As of the end of April 1983, both monkeys were responding near the 90%-correct and 250 msec response-latency criteria and are ready for electrode implants. Presentation of the two different colors will begin only after the electrodes have been implanted so that the initial responses to the novel red stimulus can be assessed.

#### H. Problems Encountered and Solutions

The surgical preparation and study of electrophysiology was delayed for two reasons. First, we were concerned that the electrode headplug would be jeopardized when we were handling monkeys that were not tamed--i.e., it might be dislodged if the monkey escaped or struggled while being removed from the cage. This was a distinct possibility with at least three monkeys, two of which turned somersaults in their cages when being retrieved, and another often escaped. Second, we felt that the electrodes might deteriorate during the long training period prior to CNV recordings.

A related problem was the extreme aggressiveness of two monkeys, precipitating attempts to tame all the monkeys by handling them a lot, giving them fruit while they were being handled, etc. The handling problems were minimized considerably by hanging short chains from the necks. This obviated the need to grasp an arm and helped immensely in removing the monkeys from their cages (most either walk or leap out), in preventing their biting (by grasping the chain closely under the chin), and in putting them in the primate chair. Because of these developments and because the monkeys were quiet once they were in the chair, extensive chair training and other taming are unnecessary. Consequently, with subsequent groups formal experimentation can be initiated much more quickly.

Motivating some of the monkeys also proved to be a problem. All except two had to be on a schedule that required obtaining their total daily fluid intake in the testing situation (when daily training was in progress, a quart of water provided Friday night was usually fully consumed by Saturday afternoon). Subsequently, we were able to test successfully just on Monday, with ad-lib watering Tuesday through Friday afternoon. Thus, after training, performance appeared to be somewhat less dependent on the state of deprivation.

During training two problems occurred. First, although the monkeys very rapidly learned to make some operant response with the right hand (in one training session if motivated), shaping to press the small bar used was too lengthy a process. A different manipulandum will be used with subsequent groups.

Shaping to maintain a hold on the ball was also difficult. One oddity is that the monkeys attributed to the ball characteristics of the bar. That is, they would move back from the bar to the ball, or pat the ball in an attempt to get reward rather than press the bar. This confusion is understandable because the ISI did have characteristics of a reinforcer (secondary). Using a manipulandum that has to be held during the ITI, rather than having separate devices to hold and press, should alleviate this problem and should also facilitate operant training.

Some difficulty was encountered in using commercially obtained software to operate the analog-to-digital converter. The software had to be modified at the level of assembly language to allow sequential sampling across channels in response to a single clock overflow flag. A hardware problem was encountered in the digital-to-analog converter when we attempted to plot waveforms. An SRI engineer traced the problem to a faulty voltage regulator, which was repaired.

The lack of an appropriate stereotaxic atlas required a more extensive effort to determine electrode coordinates than would ordinarily be required, and the placements may not be where they were aimed. A related problem is that the cynomolgus' vertical brain placement in the cranium varies greatly from one monkey to the next,

requiring the placement of depth electrodes with respect to the dura. Significant variability in the anterior-posterior plane cannot be compensated without technically demanding X-ray techniques.

Impedances were above 100 k $\Omega$  in subcortical electrodes in several monkeys and in similar electrodes placed in normal saline. However, some of the placements appear to be open and unusable, not just of high impedance. We plan to explore the use of a preamplifier with a very-high-input impedance to examine this problem further. During one phase of electrode placement, the pipette must be cut about 1 cm above the skull. This is done with a cutting wheel, and the vibration often causes saline to be extruded from the pipette, leaving bubbles. The bubbles are usually successfully removed by inserting a small wire into the pipette and adding saline. However, some bubbles could have escaped below the level of the dental cement holding them and hence remained in the pipette.

Two monkeys are being trained in a visual RT task at Stanford. For those monkeys as well, rigorous control of fluid intake was required to obtain adequate performance. Modification of computer programs and improvement of the reinforcement mechanism solved other problems encountered at Stanford.

#### I. Plans for the Coming Year

During the coming year, the monkeys now implanted will be retrained on the bar-pressing task and pairing of the WS with the IS will begin. A first task will be to determine the extent to which blinks and eye movements affect the intracerebral electrodes. This will be done by time-locking EEG samples to the occurrence of blinks. Task-related electrophysiological recordings will then begin, involving, first, attempts to obtain P300-like responses in the passive paradigm. If this is not immediately successful, we will continue with our original emphasis on the CNV, with study of the P300 in active tasks at Stanford and later at SRI by varying the proportions of WS and DS trials in the CNV paradigm.

We will record daily for two weeks to observe acquisition of the slow potentials. Once stable responses are observed, massed-unit activity will be monitored. Next, we will use systemic injections of some synaptic blocking agents (e.g., atropine) to obtain preliminary estimates of neurochemical systems involved in the CNV--in anticipation of later work using intracerebral perfusions. During this next year we will also begin working with the push-pull perfusion apparatus.

At Stanford, the monkeys now being trained in a visual RT task will be implanted with multistrand electrodes and tested in the two types of P300 paradigms described above.

J. Publications and Presentations

Rebert, C. S., and Donovan, W. J. Cortical and subcortical event-related potentials in monkeys performing the cued reaction-time task. To be presented at the Seventh International Congress on Evoked Potentials, Florence, Italy, September 1983.

Rebert, C. S. Chairman's remarks and summary. Symposium on ERPs in relation to biochemistry and pharmacology. Florence, Italy, September 1983.

K. List of Professional Personnel

Charles S. Rebert, SRI International  
William J. Donovan, SRI International  
Edward E. Davis, SRI International  
Karl H. Pribram, Stanford University  
Jeffrey E. Evans, Stanford University

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**Appendix A**

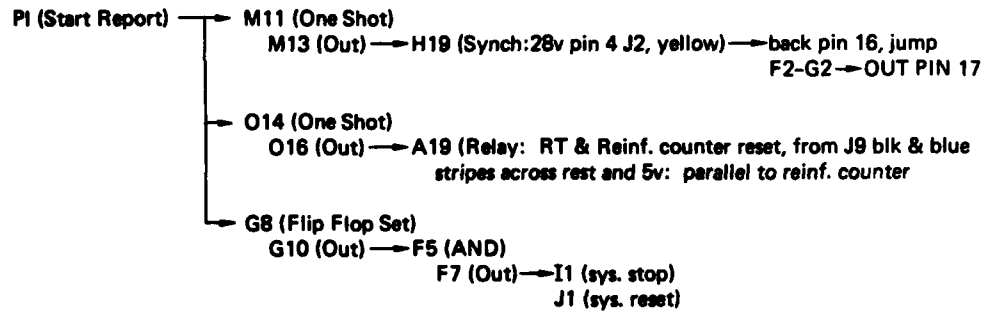
**ICONIX LOGIC SCHEMATIC**

# ICONIX CNV PROGRAM

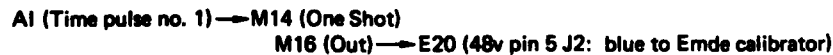
## TIMER OUTPUTS

	START Report Synch	1 CAL	2 WS	3 ABORT OFF	4 IS ON	5 IS TIMEOUT	6 IS OFF
TIME (MS)	00000	00100	00700	02690	02700	04700	14700

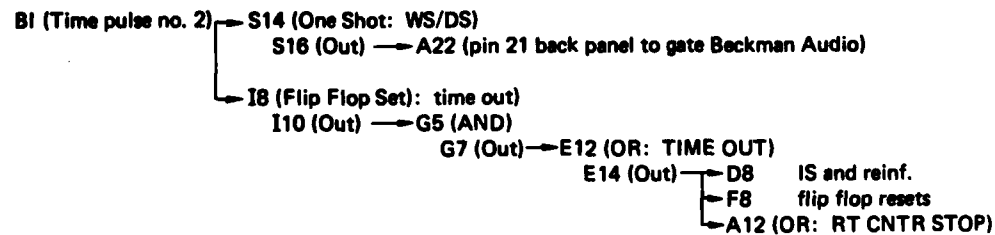
### SYNCH



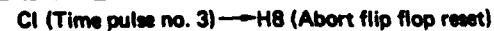
### CAL



### WS/DS & Time Out Set



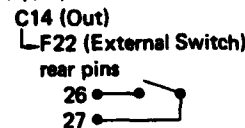
### ABORT OFF



# ICONIX CNV PROGRAM (Continued)

## IS ON/RT START

DI (Time pulse no. 4) → C11 (OR)



## REMOTE IS ON

Rear pin 17 (G2) → C12 (OR)  
from EXT. Push Button Switch

G22 (Out)

C8 (Flip Flop Set)

C10 (Out) → B19 (Relay for IS: J9

Green: orange stripe)

Q11 (One Shot)

Q13 (Out) → E19 (12 v pin 1 J2 grey → rear  
pin 31, JMP 21A&B, out  
rear pin 32: RT Start)

Connect J1-J11 for  
Normal operation  
J1-J12 for Contin. Reinf.

K22  
(front jack 11)

K21  
(front jack 1)

L22 (front jack 12)

L3 (1/4)

## IS TIME OUT

E1 (Time pulse no. 5) → G4 (AND)

E8 (Flip Flop Set: reinf. contingency)

E10 (Out) → E5 (AND)

E7 (OUT)

## RESPONSE BAR

BAR → rear pin 23

(front panel C22) → B12

Hand Switch → rear pin 25

(E22) → B11

OR

B14 (Out)

F20 (48v out pin 6 J2: white → rear pin  
28, JMP 122 & H22, out rear pin 29)

REINF CNTR

L21 (front jack 2) } For normal operation connect J2-J3;

M21 (front jack 3) } For continuous juice flow connect J3-J12 (≡)

C19 (Relay for reinf. solenoid: J9 solid blk & blue)

S11 (One Shot)

T13 (Out)

G19 (28 v pin 3 J2,  
red: total bar press CNTR)

O11 (One Shot)

J8 (Reset timeout FFlop)

O13 (Out)

E6 (AND: reinf. contingency)

F6 (AND: for abort)

A11

A12

(OR: RT Counter Stop)

A14 (Out)

Q14 (One Shot)

Q16 (Out)

F19 (12 v pin 2 J2, brown) → BERL CNTR

"STOP"

## IS OFF

F1 (Time pulse no. 6) → E11 (OR)

E14 (Out)

D8

F8

(IS & Reinf. FFlop resets)

## REMOTE IS OFF

EXT. SWITCH → rear pin 16 (F2)

OUT

E13 (OR)

E14 (Out)

**Appendix B**

**LSI-11/23 PROGRAM LISTINGS**

- CNV
- P300
- PLTCNV

```

C      PROGRAM CNV2 IS THE MASTER PROGRAM FOR RUNNING
C      THE CUED REACTION TIME STUDY, SAMPLING EIGHT
C      CHANNELS OF EITHER EVENT RELATED POTENTIALS OR
C      MULTIPLE UNIT ACTIVITY, AND ALL RELATED DATA
C      MANIPULATION AND STORAGE, VIA MENU SELECTION.
C      SOFTWARE TRIGGER INITIATES SAMPLING EVERY 10 MSEC.
C      DIGITIZED WAVEFORMS STORED IN MATRIX AS INTEGERS,
C      PRIOR TO DISPLAYING AND WRITING TO WINCHESTER DISK.
C      AT END OF SESSION, AVERAGES WRITTEN TO FLOPPY DISK; USER
C      TRANSFERS SINGLE-TRIAL DATA FROM WINCHDISK TO MAGTAPE.
C      PROGRAM CNV2
C      EXTERNAL WATCH, ITISUM
COMMON /BLOCK1/MSNEU,ISI/BLOCK2/ITIRSP/BLOCK3/IDIS/BLOCK4/MSROFF
VIRTUAL USAVG(8,400),DSAVG(8,400),YVAL(400),XVAL(400)
VIRTUAL MATRIX(8,400),TAGS(68)
REAL*8 FNAME(2)
REAL YVAL,XVAL,RLSTAT(14),USAVG,DSAVG,PREADJ,ADDEND,USCNT,DSCNT
REAL SORTOT,TOTMS
LOGICAL*1 YESNO,TASK(3),GONOW,IDMONK(8),TYPE,EXTMSG(34),COMENT(60)
LOGICAL*1 FSTRNG(14),TONE(3),NUFILE(14)
LOGICAL*1 FSPEC1(4),FSPEC2(7),FSPEC3(4),FSPECA(10),FSPECB(13),ERR1
INTEGER*2 MSOLD,MSNEU,ISI,NRESP(7),MSECT,IDIS,CALUV(8)
INTEGER*2 MATRIX,ISTAT,IBUFF(8),IBFCNT,ICHAN,TAGS,INSTAT(30)
INTEGER*2 MONTH(12),WINCHD,LCH(8),LIMAX(8),LIMIN(8),KLIPT(8)
INTEGER*4 ITIME
DATA WINCHD/3RDL1/,FNAME(1)/12RDL1FTN3 DAT/,RLSTAT/14*0.0/
DATA IBUFF/8*0/,NRESP/7*0/,INSTAT/30*0/
DATA MONTH/31,28,31,30,31,30,31,31,30,31,30,31/
DATA FSPEC1/'D','L','1',0/,FSPEC3/'D','A','T',0/
ITRIAL=0          ! Counter for # of trial
LIMIT=3770       ! Artifact window size(+ or -)
NDAYS=0
ITIRSP=0
ITISEC=0
LOOK=0           ! Automatically displayed channel
INTGR1=0
INTGR2=0
DSCNT=0          ! Zero out DS sample size
USCNT=0          ! Zero out US sample size
ISPEED=0
TYPE 8
8      FORMAT(' NAME OF MONKEY IS: ',4)
CALL GETSTR(5,IDMONK,7,ERR1) ! Input up to 7 letters of name
IF(ERR1) GO TO 4
D      CALL SCOPY(IDMONK,IDMONK,2) ! Store initial 2 letters only
CALL SCOPY('DY:XXTABL.DAT',FSTRNG) ! Initialize filename
CALL INSERT(IDMONK,FSTRNG,4,2) ! Insert 1st 2 letters of name
CALL IDATE(MON,IDAY,IYEAR) ! Access date entered upon boot
TYPE 12,MON,IDAY,IYEAR
12     FORMAT(' IS TODAY'S DATE: ',I3,'/',I3,'/',I3,' (Y or N)?',4)
ACCEPT 14,YESNO
16     FORMAT(A1)
IF(YESNO .EQ. 'Y') GO TO 24 ! If date OK, proceed
TYPE 20
20     FORMAT(' YOU FORGOT TO ENTER DATE; START ALL OVER ',/)
GO TO 832

```

```

24 OPEN(UNIT=2,NAME=FSTRNG,TYPE='OLD',DISPOSE='SAVE')
   READ (2,*) MONKEY,LMON,LDAY,LYEAR,KONDTN,KONDAY,LIMHLB,
   91STINI,ISDURA,KPCTUS,CALUV
   REWIND 2
   CLOSE(UNIT=2,DISPOSE='SAVE')
   TYPE 28,LMON,LDAY,LYEAR
28  FORMAT(/,' LAST SESSION WAS ON:',I3,'/',I3,'/',I3,/)
   LMON=MON           ! UPDATE DATE OF MOST RECENT SESSION
   LDAY=IDAY
   LYEAR=IYEAR
   KONDAY=KONDAY+1    ! UPDATE THE # OF DAYS WITHIN THIS CONDITION
   TYPE 32,KONDAY,KONDTN
32  FORMAT(/,' THIS WILL BE DAY #',I3,' OF CONDITION #',I3,' ;CHANGE? ',*)
   ACCEPT 36,YESNO
36  FORMAT(A1)
   IF(YESNO .NE. 'Y') GOTO 500
   TYPE 40
40  FORMAT(/,3X,'#1---IS-RT TESTING',/,3X,'#2---PSEUDOCONDITIONING',/,3X,
   9' #3---US-DS TESTING',/, ' WHICH # FOR NEXT CONDITION? WHAT # DAY?',
   9'(e.g.,4,1): ',*)
   ACCEPT 44,KONDTN,KONDAY
44  FORMAT(I3,I3)
   GO TO 500
48  CALL PUTSTR(7,' ENTER CHANNELS IN ASCENDING ORDER',')
52  DO 56 I=1,8
   LCH(I)=0           ! Zero out channel pointers
56  CONTINUE
   NCHS=0
   TYPE 60
60  FORMAT(' WHICH CHANNELS ARE TO BE SAMPLED(1,2,...,8/):',*)
   ACCEPT *,LCH(1),LCH(2),LCH(3),LCH(4),LCH(5),LCH(6),LCH(7),LCH(8)
   LHIGH=LCH(1)       ! Initialize high channel pointer to 1st channel
   DO 68 M=1,8
   IF(LCH(M) .EQ. 0) GOTO 72      ! Stop at empty pointer
   IF(LCH(M) .LT. LHIGH) GOTO 48  ! Only ascending order OK
   LHIGH=LCH(M)
   NCHS=NCHS+1
   TYPE 64,LCH(M)
64  FORMAT(10X,'CHANNEL',I2)
68  CONTINUE
72  TYPE 76
76  FORMAT(' IS THIS LIST CORRECT ? (Y or N):',*)
   ACCEPT 80,YESNO
80  FORMAT(A1)
   IF(YESNO .NE. 'Y') GOTO 52
   IF(MOD(IYEAR,4) .EQ. 0) MONTH(2)=29      ! If leap yr, Feb has 29
   NYEAR=MOD(IYEAR,10)                      ! Which year of the decade
   DO 84 JMON=1,MON
   NDAYS=NDAYS+MONTH(JMON)
84  CONTINUE
   NDAYS=NDAYS+IDAY-MONTH(MON)              ! Which day of the year
   IPART=(NDAYS-MOD(NDAYS,100))/100
   JPART=(MOD(NDAYS,100)-MOD(NDAYS,10))/10
   KPART=MOD(NDAYS,10)

```

```

      FSPEC2(1)=IDMONK(1)
      FSPEC2(2)=IDMONK(2)
      ENCODE (4,88,FSPEC2(3)) IPART,JPART,KPART,NYEAR
      FORMAT(4I1)
88      FSPEC2(7)=0
      CALL CONCAT(FSPEC1,FSPEC2,FSPECA,9)
      CALL CONCAT(FSPECA,FSPEC3,FSPECB,12)
      CALL IRAD50(12,FSPECB,FNAME(2))           ! Store file name
      CALL SCOPY('DL1:XXXXXX.DAT',NUFILE)
      CALL INSERT(FSPEC2,NUFILE,5,6)
      DO 96 L=1,400
      DO 92 K=1,8
      MATRIX(K,L)=0           ! Zero out single-trial matrix
      USAVG(K,L)=0           ! Zero out US average matrix
      DSAVG(K,L)=0           ! Zero out DS average matrix
92      CONTINUE
      YVAL(L)=0           ! Zero out display matrix
      XVAL(L)=0
96      CONTINUE
      DO 100 M=1,NDAYS+5
      USORDS=RAN(INTGR1,INTGR2)           ! Start RAN subrtm at new point
100     CONTINUE
      NUSRUN=0
      NDSRUN=0
      ID0="0"
      I01="4000"
      ID2="44"
      ID3="10000"
      ID5="400"

      PAUSE 'HIT RETURN TO GO'
      CALL DEVICE(-1,ID0,"164000)
      CALL IPOKE("170420,"0)           ! Turn off clock
      OPEN(UNIT=3,NAME='DL1:FTN3.DAT',FORM='UNFORMATTED',INITIALSIZE=500)
108     ITRIAL=ITRIAL+1           ! Set next trial #
      TYPE 112,ITRIAL
112     FORMAT(/,' NEXT TRIAL IS #',I3)
      IF(KPCTWS .LT. 100) GOTO 116       ! If not all US trials, skip ahead
      GO TO 124
116     IF(KPCTWS .EQ. 0) GOTO 132
      USORDS=RAN(INTGR1,INTGR2)
      CUTOFF=FLOAT(KPCTWS)/100.
      IF(USORDS .GT. CUTOFF) GO TO 128       ! If true, DS tone next
120     NDSRUN=0           ! Otherwise, US tone next
      NUSRUN=NUSRUN+1
      IF(NUSRUN .GT. ((KPCTWS/10)+3)) GOTO 128! 3 too many US tones ?
124     CALL SCOPY('US',TONE)
      IF(MONKEY .GT. 1024) GOTO 140       ! Counterbalance
      GO TO 136
128     NUSRUN=0
      NDSRUN=NDSRUN+1
      IF(NDSRUN .GT. 3) GOTO 120           ! Already had 3 DS tones ?
132     CALL SCOPY('DS',TONE)
      IF(MONKEY .GT. 1024) GOTO 136       ! Counterbalance
      GO TO 140

```



```

136 ID4="100010
    GO TO 144
140 ID4="20002
144 TYPE 148,TONE
148 FORMAT(' Next trials',3A1,'tone')
C HERE REPORT, THEN INITIALIZE OTHER STARTING VALUES
  IF(ITRIAL .EQ. 1) GOTO 168
  TYPE 152
152 FORMAT(/,' BAR-PRESSES FOR LAST TRIAL BY PHASE : ')
  DO 160 J=1,7 ! Zero out counters for bar-presSES
  TYPE 156, J,NRESP(J)
156 FORMAT(' DURING PHASE',12,' : ',13,' RESPONSES')
  NRESP(J)=0
160 CONTINUE
  MSECRT=MSECRT-2500
  TYPE 164,MSECRT
164 FORMAT(/,' REACTION TIME FOR LAST TRIAL WAS ',15,' MSEC')
168 MSECRT=17500 ! Default for when no response
  JPHASE=1 ! Reset trial phase indicator
  JDATUM=0 ! Reset pointer for storing data
  MSOLD=0 ! Counter for every msec in trial
  MSNEW=0 ! Updated msec counter
  ISI=0 ! Counter for every 10 msec
  ISYNOT=0 ! Initialize reject reason code
  MSROFF=20000 ! Initialize reinforcement timer
  IDIS="40000
  CALL DEVICE(-1,ID1,"164000)
  IF(ITRIAL .EQ. 1) GOTO 180
  CALL CCSETI(0) ! Disable ITI response counter
  CALL GTIN(ITIME)
  CALL CVTTIN(ITIME,IHRS,IMIN,ISEC,ITCK)
  JTIRSP=ITIRSP ! Store # of responses in last ITI
  ITISEC=(3600*IHRS)+(60*IMIN)+(ISEC)-ITISEC ! Msec since
  TYPE 176,ITISEC,ITIRSP ! last trial
176 FORMAT(' LAST ITI OF',14,' SEC INCLUDED',14,' BAR-PRESSES',/)
  ITIRSP=0 ! Reset ITI response tally
D NOW=0
C CLOCK WILL RUN AT 10 KHz, OVERFLOW EACH 1 MSEC
C CLOCK INTERRUPT WILL ONLY BE USED TO UPDATE MSNEW AND ISI
180 CALL CCREAD(MBPRSS,MYET,MSTILL,"164010)
  IF((MBPRSS+MYET+MSTILL).GT.0) GOTO 180
  CALL CLOCKU(1,3,-10,,WATCH,"170420,"440) ! Start clock
184 IF(MSNEW .LE. MSOLD) GOTO 184 ! Wait for next clock overflow
  MSOLD=MSNEW ! Resynch msec counters
  IF(ISI .LT. 10) GOTO 208 ! Sample or check for bar-presSES
  ISI=0
  JDATUM=JDATUM+1
  CALL AREADP(ISTAT,IBUFF,8,-7) ! Sample channel(s)
  CALL EXT812(IBUFF,8)
188 IF(ISTAT .EQ. 0) GOTO 188 ! Wait until sampling all done
  IF(ISTAT .EQ. 1) GOTO 196 ! Without any errors ?
  TYPE 192
192 FORMAT(' ERROR IN DATA COLLECTION ',/)
  CALL IPOKE("170420,"0) ! Turn off clock, then abort trial

```

```

ITRIAL=ITRIAL-1
GO TO 448      ! Terminate trial
196 DO 200 K=1,8
MATRIX(K,JDATUM)=IBUFF(K)      ! Transfer data to full matrix
200 CONTINUE
IF(MSNEW .GE. MSROFF) CALL ROFF
GO TO 184

204 ISYNOT=3      ! Experimenters vote to abort
GO TO (216,220,224,228,232,236), JPHASE ! Continue aborted trial
208 CALL CCREAD(MBPRSS,MYET,MSTILL,"164010" ! Any contact closures?
IF(MYET .GE. 4096) GOTO 204      ! Chan 13-16;Experimenter aborted
IF(MBPRSS .EQ. 0) GOTO 212      ! If none, time for next event ?
NRESP(JPHASE)=NRESP(JPHASE)+1 ! Bar-press
IF(IDIS .NE. "40200") GOTO 212 ! If DS, or abort, or phase<6
CALL REINF
IF(MSECRT .EQ. 17500) MSECRT=MSNEW      ! Note reaction time

C TRANSITION TO NEXT PHASE OF TRIAL YET ?
212 GO TO (216,220,224,228,232,236), JPHASE
216 IF(MSNEW .GT. 400) GOTO 240      ! Begin CAL trigger pulse
GO TO 184
220 IF(MSNEW .GT. 450) GOTO 244      ! End CAL trigger pulse
GO TO 184
224 IF(MSNEW .GT. 1000) GOTO 248      ! Begin tone
GO TO 184
228 IF(MSNEW .GT. 1200) GOTO 252      ! End tone
GO TO 184
232 IF(MSNEW .GT. ISON) GOTO 256      ! Turn on light
GO TO 184
236 IF(MSNEW .GT. 4000) GOTO 264      ! Stop sampling
GO TO 184
240 JPHASE=2
CALL DEVICE(-1,ID2,"164000")      ! CAL pulse ON
GO TO 184
244 JPHASE=3
CALL DEVICE(-1,ID3,"164000")      ! CAL pulse OFF
GO TO 184
248 JPHASE=4
CALL DEVICE(-1,ID4,"164000")      ! Tone ON
GO TO 184
252 JPHASE=5
CALL DEVICE(-1,ID5,"164000")      ! Tone OFF
GO TO 184
256 JPHASE=6
IF((NRESP(1)+NRESP(2)+NRESP(3)+NRESP(4)+NRESP(5)) .EQ. 0) GOTO 260
ISYNOT=2      ! Premature bar-press
260 IF(ISYNOT .GT. 0) GOTO 184
IF(SCOMP('WS',TONE) .NE. 0) GOTO 262      ! If DS trial, no IS
IDIS="40200"
262 CALL DEVICE(-1,IDIS,"164000")      ! Phase 6 events begin
GO TO 184

```

```

264 JPHASE=7          ! No more sampling but trial not over yet
    IF(ISYNOT .GT. 0) GOTO 296
    IF(IDIS .EQ. "40000) GOTO 293      ! If DS trial, skip ahead
    IDUSOK=IDIS-16384+512             ! Switch indicator lights: 6th to 7th
    CALL DEVICE(-1,IDUSOK,"164000)    ! Without altering ongoing reinf
    IDIS="1200
    IF(NRESP(6) .EQ. 0) GOTO 268      ! If no 1st bar-press yet, skip ahead
264 IF(MSROFF .EQ. 20000) GOTO 280
    GO TO 276
268 CALL CCREAD(MBPRSS,MYET,MSTILL,"164010)
    IF(MYET .GT. MSTILL) GOTO 272      ! Was there a contact closure?
    IF(MSNEW .LT. LIMRT) GOTO 268      ! More time left ?
    ISYNOT=1                          ! Too slow reaction time
    GO TO 296
272 MSECRT=MSNEW          ! Record reaction time to 1st bar-press
    CALL REINF
    NRESP(7)=1
276 IF(MSNEW .LT. MSROFF) GOTO 276      ! Wait for reinforcement delivery
    CALL ROFF
    GO TO 280
278 IF(IDIS .NE. "1200) GOTO 278
280 CALL CCREAD(MBPRSS,MYET,MSTILL,"164010)
    IF(MBPRSS .EQ. 0) GO TO 284        ! If no bar-press, skip ahead
    NRESP(7)=NRESP(7)+1
    CALL REINF
284 IF(MSROFF .EQ. 20000) GOTO 292
288 IF(MSNEW .LE. MSROFF) GOTO 288      ! Wait for reinf relay offset
    CALL ROFF
292 IF(MSNEW .GE. ISOFF) GOTO 296      ! End of trial yet ?
    GO TO 280

293 IDIS="1000
    CALL DEVICE(-1,IDIS,"164000)      ! Phase M7 light on
294 CALL CCREAD(MBPRSS,MYET,MSTILL,"164010)
    IF(MBPRSS .EQ. 0) GOTO 295
    NRESP(7)=NRESP(7)+1
295 IF(MSNEW .LT. ISOFF) GOTO 294

296 CALL DEVICE(-1,IDO,"164000)        ! Light OFF, end reinf
    CALL IPOKE("170420,"0)           ! Turn off clock during ITI
    CALL GTIN(ITIME)
    CALL CUTTIN(ITIME,IHRS,IMIN,ISEC,ITCK)
    ITISEC=(3600*IHRS)+(60*IMIN)+(ISEC)
    NOW=ITISEC                        ! Store time at end of this trial
C   BEGIN INTER-TRIAL INTERVAL RESPONSE-COUNTING,
C   THEN REVIEW DATA FOR ARTIFACTS, THEN STORE RAW DATA
C   TEMPORARILY ON WINCH DISK AS A SINGLE RECORD PER TRIAL.
    CALL CCSETI(ITISUM,"164010,"270)  ! Count # ITI bar-presses
    CALL WINDOW(MATRIX,LHIGH,LIMIT,LCH,LIMAX,LIMIN,KLIPT)
    TYPE 300
300 FORMAT(/, ' ENTER COMMENT(up to 59 chars) FOR THIS TRIAL:',/)
    CALL GETSTR(5,COMENT,59)
    CALL STRPAD(COMENT,59)

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```

TAGS(1)=MONKEY
TAGS(2)=MON
TAGS(3)=IDAY
TAGS(4)=IYEAR
TAGS(5)=KONDTN
TAGS(6)=KONDAY
TAGS(7)=KPCTUS
TAGS(8)=LIMHLD
TAGS(9)=ISTIMI
TAGS(10)=ISDURA
TAGS(11)=LIMIT
TAGS(12)=NCHS
DO 304 I=13,20
TAGS(I)=LCH(I-12)
304 CONTINUE
TAGS(21)=USCNT
TAGS(22)=DSCNT
TAGS(23)=NSECRT-2500
TAGS(24)=ID4
TAGS(25)=JTIRSP
TAGS(26)=NOW
TAGS(27)=ITRIAL
TAGS(28)=ISYNOT
DO 308 I=29,35
TAGS(I)=NRESP(I-28)
308 CONTINUE
DO 312 J=36,43
TAGS(J)=LIMAX(J-35)
TAGS(J+8)=LIMIN(J-43)
TAGS(J+16)=KLIPT(J-51)
TAGS(J+24)=CALUV(J-59)
312 CONTINUE
WRITE (3) TONE,TAGS,((MATRIX(LCH(N),K),K=1,400),M=1,NCHS),CONENT
IF(ISYNOT .GT. 0) GOTO 324
DO 320 I=1,8
IF(KLIPT(LCH(I)) .GT. 5) ISYNOT=4      ! More than 5 artifacts?
IF(LCH(I) .EQ. LHIGH) GOTO 324
320 CONTINUE
324 GO TO (328,364,372,380,388),(ISYNOT+1)

328 IF(SCOMP('DS',TONE) .EQ. 0) GOTO 348      ! If DS trial,update DSAVG
DO 340 L=1,8
MCH=LCH(L)
DO 332 N=1,400
PREADJ=USAVG(MCH,N)
ADDEND=MATRIX(MCH,N)
USAVG(MCH,N)=(PREADJ+USCNT+ADDEND)/(USCNT+1.0)
332 CONTINUE
336 IF(MCH .EQ. LHIGH) GOTO 344
340 CONTINUE
344 USCNT=USCNT+1.0
GO TO 396

```

```

348      DO 354 L=1,8
          MCH=LCH(L)
          DO 352 N=1,400
              PREADJ=DSAVB(MCH,N)
              ADDEND=MATRIX(MCH,N)
              DSAVB(MCH,N)=(PREADJ+DSCNT+ADDEND)/(DSCNT+1.0)
352      CONTINUE
          IF(MCH.EQ. LHIGH) GOTO 360
354      CONTINUE
360      DSCNT=DSCNT+1.0
          GO TO 396

364      TYPE 368
368      FORMAT(/,' TOO SLOW REACTION TIME; DATA NOT ADDED TO AVERAGE')
          PAUSE 'HIT RETURN'
          GO TO 396
372      TYPE 376
376      FORMAT(/,' PREMATURE BAR-PRESS; DATA NOT ADDED TO AVERAGE')
          PAUSE 'HIT RETURN'
          GO TO 396
380      TYPE 384
384      FORMAT(/,' EXPERIMENTER INTERVENED; DATA NOT ADDED TO AVERAGE')
          PAUSE 'HIT RETURN'
          GO TO 396
388      TYPE 392
392      FORMAT(/,' TOO MANY ARTIFACTS; DATA NOT ADDED TO AVERAGE')
          PAUSE 'HIT RETURN'

396      IF(ISPEED.EQ. 1) GOTO 448
          IDCH=LOOK
          DO 404 N=1,400
              YVAL(N)=MATRIX(IDCH+1,N)+380
              XVAL(N)=N+150+N
          CONTINUE
          CALL GINIT(1)           ! Set default parameters
          CALL GEND              ! Go to transparent mode
          CALL STXOX
          CALL STOTU(7)          ! Graphic output on console
          CALL STERR(2)          ! Print error and warning messages
          TYPE 412
          FORMAT(25(/))         ! Clear screen
          DO 416 N=1,400
              CALL POINT(XVAL(N),YVAL(N)) ! Plot values
          CONTINUE
          CALL GEND              ! Go to transparent mode
          CALL HOLD              ! Wait for carriage return
          CALL ERASE             ! Clear screen
          CALL GEND              ! Go to transparent mode
          TYPE 420
          FORMAT(' VIEW MORE SINGLE-TRIAL DATA (Y or N) ? ',0)
          ACCEPT 424,YESNO
          FORMAT(A1)
          IF(YESNO.EQ. 'Y') GOTO 464

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428      TYPE 428
428      FORMAT(' VIEW MORE AVERAGE DATA (Y or N) ? ',*)
      ACCEPT 432,YESNO
432      FORMAT(A1)
      IF(YESNO.EQ.'Y') GO TO 476
D       GO TO 448
D436     TYPE 440
D440     FORMAT(/,' Enter "G" to start next trial, "M" for menu',*)
D       ACCEPT 444,YESNO
D444     FORMAT(A1)
D       IF(YESNO.EQ.'G') GO TO 108

448      TYPE 450
450      FORMAT(/,' HIT RETURN FOR NEXT TRIAL, OR: ')
      TYPE 452
452      FORMAT(' PICK FROM MENU : ??,QT,EX,DS,DA,CP,PL,...',*)
      ACCEPT 456,TASK(1),TASK(2)
456      FORMAT(2A1)
      IF(SCOMP('??',TASK).EQ.0) GO TO 458      ! Display menu
      IF(SCOMP('QT',TASK).EQ.0) GO TO 456      ! Quit session
      IF(SCOMP('EX',TASK).EQ.0) GO TO 460      ! Exit session
      IF(SCOMP('DS',TASK).EQ.0) GO TO 464      ! Display data
      IF(SCOMP('DA',TASK).EQ.0) GO TO 476      ! Display average
      IF(SCOMP('CP',TASK).EQ.0) GO TO 500      ! Change parameters
      IF(SCOMP('PL',TASK).EQ.0) GO TO 648      ! Plot data
      GO TO 108

458      TYPE 460
460      FORMAT(' MENU IS AS FOLLOWS: ??---EXPLAIN MENU',/,23X,
7'QT---QUIT SESSION(DELETE DATAFILES)',/,23X,'EX---EXIT ',
7'FROM SESSION(SAVE DATAFILES)',/,23X,'DS---DISPLAY SINGLE
7'TRIAL DATA',/,23X,'DA---DISPLAY AVERAGES',/,23X,
7'CP---CHANGE PARAMETERS',/,23X,'PL---PLOT DATA',/,/)
      GO TO 448

464      TYPE 468
468      FORMAT(' VIEW DATA FROM WHICH CHANNEL (1-8) ? ',*)
      ACCEPT 472,IDCH
472      FORMAT(I3)
      IDCH=IDCH-1
      IF(IDCH.LT.0 .OR. IDCH.GT.7) GO TO 464
      GO TO 400

476      TYPE 480
480      FORMAT(' US OR DS ? WHICH CHANNEL ? (e.g. U4 or U8 or D1)',*)
      ACCEPT 484,TYPE,NUMCH
484      FORMAT(A1,I1)
      IF(TYPE.NE.'U' .AND. TYPE.NE.'D') GO TO 476
      IF(NUMCH.LT.1 .OR. NUMCH.GT.8) GO TO 476
      IF(TYPE.EQ.'D') GO TO 492
      DO 488 K=1,400
      YVAL(K)=USAVG(NUMCH,K)+380.
      XVAL(K)=K+150+K
488      CONTINUE
      GO TO 408

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492      DO 496 K=1,400
        YVAL(K)=DBAUG(NUMCH,K)+380.
        XVAL(K)=K+150+K
496      CONTINUE
        GO TO 408
500      TYPE 504,KPCTUS,LINHLD,LIMIT,(LOOK+1)
504      FORMAT(/,' PARAMETERS ARE AS FOLLOWS:',/, ' 1) X OF US TRIALS= ',I4,
        9/, ' 2) LIMITED HOLD(msec)= ',I5,/, ' 3) ARTIFACT LIMIT= +/- ',I6,/,
        9' 4) CHANNEL DISPLAYED AT END OF EACH TRIAL= ',I2,/, ' 5) CAL PULSES:',)
        DO 512 M=1,8
          TYPE 508,M,CALUV(M)
508      FORMAT(3X,'CH',I2,': ',I5,'uV')
512      CONTINUE
          TYPE 516,ISTIMI,ISDURA,ISPEED
516      FORMAT(/,' 6) INTER-STIMULUS INTERVAL(210-2990 msec)= ',I5,/,
        9' 7) IS DURATION(>2800 msec)= ',I6,/, ' 8) SPEED(0=slow;1=fast)= ',I2)
          TYPE 520
520      FORMAT(' ENTER NUMBER OF ITEM TO BE CHANGED, OR 0 IF NONE : ',)
          ACCEPT 524,NPARAM
524      FORMAT(I2)
          GO TO (644,528,540,552,564,596,608,620,632), (NPARAM+1)

528      TYPE 532
532      FORMAT(' WHAT X(0-100) TRIALS SHOULD BE US ? ',)
          ACCEPT 536,KPCTUS
536      FORMAT(I4)
          GO TO 500
540      TYPE 544
544      FORMAT(' HOW MANY MSEC FOR LIMITED HOLD ? ',)
          ACCEPT 548,LINHLD
548      FORMAT(I5)
          GO TO 500
552      TYPE 556
556      FORMAT(' NEW WINDOW LIMIT= ',)
          ACCEPT 560,LIMIT
560      FORMAT(I6)
          GO TO 500
564      TYPE 568
568      FORMAT(' CHANNEL TO LOOK AT FIRST(1-8) = ',)
          ACCEPT 572,LOOK
572      FORMAT(I2)
          LOOK=LOOK-1
          GO TO 500
576      DO 584 J=1,8
          TYPE 580,J,CALUV(J)
580      FORMAT(3X,'CH ',I1,' CAL= ',I5,'uV')
584      CONTINUE
          TYPE 588
588      FORMAT(/,' CHANGE CAL uV VALUES ? ',)
          ACCEPT 592,YESNO
592      FORMAT(A1)
          IF(YESNO .NE. 'Y') GO TO 500
596      TYPE 600
600      FORMAT(' ENTER CHANNEL N, THEN NEW uV CALIBRATION, e.g. 5.75 : ',)

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        ACCEPT 604,K,CALUV(K)
604    FORMAT(I2,I4)
        GO TO 576
608    TYPE 612
612    FORMAT(/,' ENTER NEW INTER-STIMULUS INTERVAL(210-2990 msec): ',%)
        ACCEPT 616,ISTIMI
616    FORMAT(I5)
        IF(MOD(ISTIMI,10) .EQ. 0) GOTO 500
        CALL PUTSTR(7,' MULTIPLES OF 10 ONLY',' ')
        GO TO 608
620    TYPE 624
624    FORMAT(/,' ENTER NEW IS DURATION(>2800 msec): ',%)
        ACCEPT 628,ISDURA
628    FORMAT(I6)
        IF(MOD(ISDURA,10) .EQ. 0) GOTO 500
        CALL PUTSTR(7,' MULTIPLES OF 10 ONLY',' ')
        GO TO 620
632    TYPE 636
636    FORMAT(/,' Enter "1" for fast, or "0" for slow mode: ',%)
        ACCEPT 640,ISPEED
640    FORMAT(I2)
        GO TO 448
644    ISON=ISTIMI+1000      ! Compute when to turn light on
        ISOFF=ISON+ISDURA ! Compute when to shut light off
        LIMRT=ISON+LIMHLD  ! Compute limited hold cutoff(msec)
        IF(ITRIAL .EQ. 0) GOTO 52
        GO TO 448

648    TYPE 652
652    FORMAT(' THIS OPTION NOT YET AVAILABLE ',/)
        GO TO 448
656    CLOSE(UNIT=3,DISPOSE='DELETE')
        CALL IPOKE("I70420","0")      ! Shut off clock
        CALL CCSETI(0)
        GO TO 832
660    TYPE 664
664    FORMAT(' OK TO END THIS SESSION (Y or N): ?',%)
        ACCEPT 668,YESNO
668    FORMAT(A1)
        IF(YESNO .NE. 'Y') GOTO 448
        CLOSE(UNIT=3,DISPOSE='SAVE')   ! Save single-trial data
        ICHANL=IGETC()
        IF(ICHANL .LT. 0) STOP 'NO CHANNEL'
        IF(IFETCH(WINCHD) .NE. 0) STOP 'FATAL ERROR FETCHING HANDLER'
        CALL IRENAH(ICHANL,FNAME)
        CALL ICLOSE(ICHANL)
        CALL IFREEC(ICHANL)
        CALL IPOKE("I70420","0")      ! Shut off clock
        CALL CCSETI(0)                ! Ignore contact closures
C      An "ok" trials: neither aborted, nor rejected due to artifacts
        TOTMS=0      ! Sum RTs for ok WS trials
        SQRTOT=0     ! Sum squared RTs
        NREINF=0     ! Total # reinforcements
        DO 672 J=1,8
        KLIPT(J)=0   ! Total # artifacts on art.-rej. WS trials
672    CONTINUE

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D      PAUSE / UPDATE TABLE/
      OPEN(UNIT=2,NAME=FSTRNG,TYPE='OLD',DISPOSE='SAVE')
      WRITE (2,*) MONKEY,LMON,LDAY,LYEAR,KONDTN,KONDAY,LINHLD,
      9ISTINI,ISDURA,KPCTUS,CALUV
      CLOSE(UNIT=2,DISPOSE='SAVE')

D      PAUSE / NOW REOPEN RENAMED FILE/
D      OPEN(UNIT=3,ERR=725,NAME=NUFILE,FORM='UNFORMATTED',TYPE='OLD')
      OPEN(UNIT=3,ERR=725,NAME=NUFILE,READONLY,TYPE='OLD',
      7FORM='UNFORMATTED',DISPOSE='SAVE')
D      PAUSE / NOW REVIEW RECORDS/
      DO 724 NREC=1,ITRIAL
      READ (3) TONE,TAGS,((MATRIX(M,K),K=1,400),M=1,NCHS),COMENT
      NBP5=0 / # of bar-presses in phases 1-5
      IF(TAGS(27) .GT. 1) GOTO 684
      DO 680 I=1,20
      INSTAT(I)=TAGS(I) / Session-wide parameters
680      CONTINUE
      INSTAT(21)=USCNT
      INSTAT(22)=DSCNT
684      INSTAT(23)=INSTAT(23)+1 / # of trials
      INSTAT(30)=INSTAT(30)+TAGS(25) / # III bar-presses
      DO 688 L=29,33
      NBP5=NBP5+TAGS(L) / # phase 1-5 bar-presses
688      CONTINUE
      NBPIS=TAGS(34)+TAGS(35) / # IS bar-presses
      INSTAT(29)=INSTAT(29)+NBP5+NBPIS+TAGS(25) / All bar-presses
      IF(SCOMP('DS',TONE) .EQ. 0) GOTO 716 / If DS, skip ahead
      INSTAT(24)=INSTAT(24)+1 / # of US trials
      IF(TAGS(28) .GT. 0) GOTO 692 / If not ok trial, skip ahead
      TOTHS=TOTHS+TAGS(23) / RT tally for ok US trials
      SQRTOT=SQRTOT+FLOAT(TAGS(23))*2 / RT squared tally
      NREINF=NREINF+NBPIS / Reinforcement tally

692      GO TO (724,696,700,704,708),(TAGS(28)+1) / ISYNOT value?
696      INSTAT(25)=INSTAT(25)+1 / US reject--too slow RT
      GO TO 724
700      INSTAT(26)=INSTAT(26)+1 / US reject--premature bar-press
      GO TO 724
704      INSTAT(27)=INSTAT(27)+1 / US reject--E opted to abort
      GO TO 724
708      INSTAT(28)=INSTAT(28)+1 / US reject--too many artifacts
      DO 712 J=1,8
      KLIPT(J)=KLIPT(J)+TAGS(J+51) / Artifact tally:US reject;not aborted
712      CONTINUE
      GO TO 724
716      IF(ISYNOT .GT. 1) GOTO 724 / Skip DS trials with problems
720      RLSTAT(6)=RLSTAT(6)+NBPIS / DS trial bar-presses:phases 6+7
      GO TO 724
724      CONTINUE
      CLOSE(UNIT=3,DISPOSE='SAVE')
      GO TO 727
725      TYPE 726
726      FORMAT(' ERROR REOPENING FILE ON DL1 ')
      GO TO 832

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D      PAUSE ' COMMENT HEADER'
727    PRINT 728
728    FORMAT(5(/),15X,'LISTING OF SESSION COMMENTS:',/)
D      PAUSE ' REOPEN DL1 FILE'
      OPEN(UNIT=3,NAME=NUFILE,FORM='UNFORMATTED',TYPE='OLD')
732    READ(3,END=740) TONE,TAGS,((MATRIX(LCH(M),K),K=1,400),M=1,NCHS),COMENT
      PRINT 736,TAGS(27),(COMENT(K),K=1,LEN(COMENT))
736    FORMAT(' TRIAL #',I3,' : ',59(A1))
      GO TO 732
740    CLOSE(UNIT=3,DISPOSE='SAVE')
D      PAUSE ' COMPUTE STATS'
      X1=INSTAT(24) ! Total # of US trials
      X2=INSTAT(25) ! # of US aborts due to slow RT
      X3=INSTAT(26) ! " due to premature response
      X4=INSTAT(27) ! " due to E intervention
      X5=INSTAT(28) ! # of US rejects due to artifacts
      X6=X1-X2-X3-X4-X5 ! X6 (=USCNT) : # of ok US trials
      IF(X1 .LE. 0) GOTO 752
      RLSTAT(1)=((X2+X3+X4)/X1)*100. ! % of US trials aborted
      RLSTAT(2)=(X5/X1)*100. ! % of US trials artifact-rejected
      IF(X6 .LE. 0.) GOTO 744 ! If no US trials OK, skip ahead
      RLSTAT(3)=TOTMS/X6 ! Mean RT for ok US trials
      RLSTAT(4)=SQRT(SQRTOT/X6-(RLSTAT(3)**2)) ! Stand. dev.
      RLSTAT(5)=FLOAT(NREINF)/X6 ! Avg # of reinf per ok US trial
744    IF(X5 .LE. 0.) GOTO 752 ! If no US artifact-rejects, skip ahead
      DO 748 I=1,8
      RLSTAT(I+6)=FLOAT(KLIPT(I))/X5 ! Avg # artifacts per US reject
748    CONTINUE
752    IF(DSCNT .LE. 0.) GOTO 756
      RLSTAT(6)=RLSTAT(6)/DSCNT ! Avg # resp(phases 6+7) for ok DS
D      PAUSE ' PRINT STATS'
756    PRINT 760
760    FORMAT(5(/))
      CALL PUTSTR(6,IDMONK,' ')
      PRINT 764,(INSTAT(1),I=1,4)
764    FORMAT(/,' Monkey #',I5,5X,' Dates:',I3,'/',I3,'/',I3)
      PRINT 768,(INSTAT(J),J=5,7)
768    FORMAT(/,' Condition #',I2,' ; Day #',I3,' ; I US trials=',I3)
      PRINT 772,(INSTAT(K),K=8,10)
772    FORMAT(/,' Limited hold(ms)=' ,I5,/, ' Inter-stimulus interval(ms)=' ,
9I5,/, ' IS duration(ms)=' ,I6)
      PRINT 776,(INSTAT(L),L=11,(INSTAT(12)+12))
776    FORMAT(/,' Artifact limit= +/-',I6,/,/, ' Sampled the following',I2,
9' channels:',8(IX,I2))
      PRINT 780,INSTAT(21),INSTAT(22)
780    FORMAT(/,' Sample size for US average=',I3,' ; for DS average=',I3)
      PRINT 784,INSTAT(23),INSTAT(24)
784    FORMAT(/,' Total # trials=',I3,/, ' # of US trials=',I3)
      PRINT 788,(INSTAT(N),N=25,28)
788    FORMAT(/,' # aborts due to slow RT=',I3,/, ' # aborts due to premature
9 bar-pressing=',I3,/, ' # aborts due to E opting to abort=',I3,/,
9' # rejects due to artifacts=',I3)
      PRINT 792,INSTAT(29),INSTAT(30)
792    FORMAT(/,' Total # bar-presses=',I5,/, ' All bar-presses during
9 ITIs=',I5)

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PRINT 796,RLSTAT(1),RLSTAT(2)
796  FORMAT(/,' Z US trials aborted=',F4.0,/, ' Z of US trials rejected
9 due to artifacts=',F4.0)
PRINT 800,RLSTAT(3),RLSTAT(4)
800  FORMAT(/,' Mean RT(ms) for "ok" US trials=',F7.1,/,
9' Standard deviation for these RTs=',F10.4)
PRINT 804,RLSTAT(5),RLSTAT(6)
804  FORMAT(/,' For ok US trials,mean # reinforcements=',F6.3,/,
9' For ok DS trials,mean # bar-presses(in phases 6+7)=',F6.3)
PRINT 808,(RLSTAT(K),K=7,14)
808  FORMAT(/,' For artifact-rejected US trials,mean # deviant points:',
9/,2X,'Chan 1',2X,'Chan 2',2X,'Chan 3',2X,'Chan 4',2X,'Chan 5',
92X,'Chan 6',2X,'Chan 7',2X,'Chan 8',/,8(2X,F6.3))

D  PAUSE  ' CREATE BY FILE'
CALL INSERT(FSPEC2,FSTRNG,4,6)
OPEN(UNIT=2,NAME=FSTRNG,TYPE='NEW',FORM='UNFORMATTED',DISPOSE='SAVE')
WRITE (2) (INSTAT(N),N=1,30),(RLSTAT(N),N=1,14)
DO 816 I=1,NCHS
DO 812 J=1,400
YVAL(J)=USAVG(LCH(I),J)      ! MULT BY USCNT?
812  CONTINUE
WRITE (2) (YVAL(N),N=1,400)
816  CONTINUE
DO 824 K=1,NCHS
DO 820 L=1,400
YVAL(L)=DSAVG(LCH(K),L)      ! MULT BY DSCNT?
820  CONTINUE
WRITE (2) (YVAL(N),N=1,400)
824  CONTINUE
CLOSE(UNIT=2,DISPOSE='SAVE')

TYPE 828
828  FORMAT(/,' SESSION IS OVER; PUT NEW FILE ON MAGTAPE BY TYFING:',/)
CALL SCOPY('COPY DL1:XXXXXX.DAT HT:XXXXXX.DAT ',EXTMSG)
CALL INSERT(FSPEC2,EXTMSG,10,6)
CALL INSERT(FSPEC2,EXTMSG,24,6)
CALL PUTSTR(7,EXTMSG,' ')
832  CALL EXIT
END

SUBROUTINE WATCH
COMMON /BLOCK1/MSNEU,ISI
MSNEU=MSNEU+1      ! Update msec counter
ISI=ISI+1          ! Update inter-sample interval timer
CALL IPOKE("170420,"133) ! Clear overflow flag
RETURN
END

```

```

SUBROUTINE ITISUM
COMMON /BLOCK2/ITIRSP
836 CALL CCREAD(MBPRSS,MYET,MSTILL,"164010)
IF(MSTILL .GT. 0) GOTO 836
ITIRSP=ITIRSP+1      ! Update M of ITI bar-presses
CALL IPOKE("164010,"100)
RETURN
END

SUBROUTINE WINDOW(MATRIX,LHIGH,LIMIT,LCH,LIMAX,LIMIN,KLIPT)
VIRTUAL MATRIX(8,400)
INTEGER*2 MATRIX,LCH(8),LIMAX(8),LIMIN(8),KLIPT(8)
DO 844 L=1,8
IF(LCH(L) .EQ. 0) GOTO 848      ! End at 1st null pointer
NX=LCH(L)      ! Point to next channel with data
LINTOP=MATRIX(NX,1)      ! Initial max value
LIMBOT=MATRIX(NX,1)      ! Initial min value
KLIPT(NX)=0      ! Reset artifact counter
DO 840 M=1,400
IF(IABS(MATRIX(NX,M)) .GT. LIMIT) KLIPT(NX)=KLIPT(NX)+1
LINTOP=MAX0(MATRIX(NX,M),LINTOP)
LIMBOT=MIN0(MATRIX(NX,M),LIMBOT)
840 CONTINUE
LIMAX(NX)=LINTOP      ! Store highest datum for Chan n
LIMIN(NX)=LIMBOT      ! Store lowest datum for Chan n
844 CONTINUE
848 RETURN
END

SUBROUTINE REINF
COMMON/BLOCK1/MSNEW,ISI/BLOCK3/IDIS/BLOCK4/MSROFF
INTEGER*2 IDIS
IDTANG=IDIS+1024      ! Set bit #10 of HCO register
CALL DEVICE(-1,IDTANG,"164000)
MSROFF=MSNEW + 50      ! Set relay operation time
RETURN
END

SUBROUTINE ROFF
COMMON/BLOCK3/IDIS/BLOCK4/MSROFF
INTEGER*2 IDIS
CALL DEVICE(-1,IDIS,"164000)      ! Clear bit #10 of HCO register
MSROFF=20000
RETURN
END

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C      PROGRAM P300 IS THE MASTER PROGRAM FOR RUNNING
C      THE UNCUELED TONE PROTOCOL, SAMPLING FOUR
C      CHANNELS OF EITHER EVENT RELATED POTENTIALS OR
C      MULTIPLE UNIT ACTIVITY, AND ALL RELATED DATA
C      MANIPULATION AND STORAGE, VIA MENU SELECTION.
C      SOFTWARE TRIGGER INITIATES SAMPLING EVERY X MSEC.
C      DIGITIZED WAVEFORMS STORED IN MATRIX AS INTEGERS,
C      ARTIFACT-REJECTED AND, OPTIONALLY, DISPLAYED.
C      AT END OF EACH SET, AVERAGES WRITTEN TO FLOPPY DISK.
C      *****C=common ; Higher probability*****
C      *****R=rare ; Lower probability*****

PROGRAM P300
EXTERNAL WATCH
COMMON /BLOCK1/MATRIX,JDATUM,IBUFF
VIRTUAL HISUM(4,400),LOSUM(4,400),YVAL(400),XVAL(400)
REAL YVAL,XVAL,HISUM,LOSUM,HICNT,LOCNT
LOGICAL*1 YESNO,TASK(3),IDSUBJ(8),TYPE,ICHAR
LOGICAL*1 FSTRNG(14),TONE(4),FSPEC(7),ERR1,ERR2,IDHIHZ
INTEGER*2 ISTAT,IBUFF(4),IBFCNT,ICHAN,INSTAT(30),CALUV(4)
INTEGER*2 MONTH(12),LEVEL(4),KLIPT(4),MATRIX(4,400)
INTEGER*4 ITIME
DATA IBUFF/4*0/,INSTAT/30*0/,CALUV/4*25/
DATA MONTH/31,28,31,30,31,30,31,31,30,31,30,31/
DATA LEVEL/4*0/,KLIPT/4*0/

LIMIT=4000          ! Artifact window size(+ or -)
NDAYS=0
ITISEC=0
LOOK=0              ! Automatically displayed channel
INTGR1=0
INTGR2=0
ISPEED=1            ! Skip single-trial display
IEPOKI=0             ! 0 sec between epochs
NTIX=25              ! Sample every 2.5 msec
IEPOCH=1000          ! Epoch lasting 1000 msec
ITIX=-25             ! Clock overflow every 2.5 msec
NTRIAL=1             ! Total # trials
ID0="0"              ! Bit pattern during ITI
ID1="4000"           ! " " " phase 1
ID2="44"             ! " " " " 2
ID3="10000"          ! " " " " 3
ID5="400"            ! " " " " 5
NSET=0               ! Initialize counter for data sets
KNIPCT=80            ! Initialize % of common tones

4      TYPE B
8      FORMAT(' NAME OF SUBJECT IS: ',*)
      CALL GETSTR(5,IDSUBJ,7,ERR1) ! Input up to 7 letters of name
      IF(ERR1) GOTO 4
      CALL SCOPY('DY:XXXXXX.DAT',FSTRNG) ! Initialize filename

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CALL IDATE(MON,IDAY,IYEAR)      ! Access date entered upon boot
TYPE 12,MON,IDAY,IYEAR
12  FORMAT(' IS TODAY'S DATE:',I3,'/',I3,'/',I3,' (Y or N)?',%)
ACCEPT 14,YESNO
14  FORMAT(A1)
    IF(YESNO.EQ.'Y') GOTO 22      ! If date OK, proceed
    TYPE 20
20  FORMAT(' YOU FORGOT TO ENTER DATE; START ALL OVER ',/)
    GO TO 832

22  TYPE 24
24  FORMAT(/,' ENTER NEW FILE NAME (up to 6 letters/numbers) : ',%)
    CALL GETSTR(5,FSPEC,6,ERR2)
    IF(ERR2) GOTO 22
    FSPEC(7)=0
    TYPE 26,(FSPEC(J),J=1,6)
26  FORMAT(/,2X,6A1,'.DAT WILL BE WRITTEN ONTO FLOPPY DISK')

    NCHS=4
    TYPE 38
38  FORMAT(/,' FOUR CHANNELS ARE TO BE SAMPLED: 1,2,3, and 4')

    IF(MOD(IYEAR,4).EQ.0) MONTH(2)=29      ! If leap yr, Feb has 29
    NYEAR=MOD(IYEAR,10)                    ! Which year of the decade?
    DO 40 JMON=1,MON
    NDAYS=NDAYS+MONTH(JMON)
40  CONTINUE
    NDAYS=NDAYS+IDAY-MONTH(MON)            ! Which day of the year

    DO 44 M=1,NDAYS+5
    RANNUH=F*N(INTOR1,INTOR2)              ! Start RAN subrtm at new point
44  CONTINUE

C    *****OPEN DATA FILE FOR THIS SESSION*****
    CALL INSERT(FSPEC,FSTRNG,4,6)          ! Open session's data file
    OPEN(UNIT=2,NAME=FSTRNG,TYPE='NEW',FORM='UNFORMATTED',DISPOSE='SAVE')

C    *****RESET PARAMETERS*****
50  NSET=NSET+1
51  TYPE 52,KHIPCT,LIMIT,(LOOK+1)
52  FORMAT(/,' PARAMETERS ARE AS FOLLOWS:',/, ' 1) % OF COMMON TONES
    9 = ',I4,/, ' 2)ARTIFACT LIMIT= +/- ',I6,/,
    9' 3)CHANNEL DISPLAYED AT END OF EACH TRIAL= ',I2,/, ' 4)CAL PULSES:')
    DO 54 M=1,4
    TYPE 53,M,CALUV(M)
53  FORMAT(3X,'CH',I2,' : ',I5,'uV')
54  CONTINUE
    TYPE 55,(NTIX+40),IEPOK1,ISPEED,NSET
55  FORMAT(/,' 5)EPOCH DURATION = ',I5,' msec',/, ' 6)Inter-
    9epoch interval = ',I6,' sec',/, ' 7)SPEED(0=slow;1=fast)= ',I2,
    9/, ' 8) # OF NEXT DATA SET = ',I4)
    TYPE 56
56  FORMAT(' ENTER NUMBER OF ITEM TO BE CHANGED, OR 0 IF NONE : ',%)
    ACCEPT 57,NPARAM
57  FORMAT(I2)

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58      GO TO (90,58,61,64,67,75,78,81,86), (NPARAM+1)
59      TYPE 59
60      FORMAT(' WHAT % OF OCCURRENCE (50< % <100) FOR COMMON TONES ? ',%)
        ACCEPT 60,KHIPCT
        FORMAT(I4)
        IF(KHIPCT .LT. 50) GOTO 58
        GO TO 51

61      TYPE 62
62      FORMAT(' NEW WINDOW LIMIT= ',%)
        ACCEPT 63,LIMIT
63      FORMAT(I6)
        GO TO 51

64      TYPE 65
65      FORMAT(' CHANNEL TO LOOK AT FIRST(1-4) = ',%)
        ACCEPT 66,LOOK
66      FORMAT(I2)
        LOOK=LOOK-1
        GO TO 51

67      DO 69 J=1,4
        TYPE 68,J,CALUV(J)
68      FORMAT(3X,'CH ',I1,' CAL= ',I5,'uV')
69      CONTINUE
        TYPE 70
70      FORMAT(/,' CHANGE CAL uV VALUES ? ',%)
        ACCEPT 71,YESNO
71      FORMAT(A1)
        IF(YESNO .NE. 'Y') GOTO 51

72      TYPE 73
73      FORMAT(' ENTER CHANNEL #, THEN NEW uV CALIBRATION, e.g. 3,75 : ',%)
        ACCEPT 74,K,CALUV(K)
74      FORMAT(I2,I4)
        GO TO 67

75      TYPE 76
76      FORMAT(/,' ENTER NEW EPOCH DURATION (msec): ',%)
        ACCEPT 77,IEPOCH
77      FORMAT(I5)
        NTIX=IEPOCH/(40)
        ITIX=-1*NTIX
        GO TO 51

78      TYPE 79
79      FORMAT(/,' ENTER NEW INTER-EPOCH INTERVAL (sec): ',%)
        ACCEPT 80,IEPOKI
80      FORMAT(I6)
        GO TO 51

81      TYPE 82
82      FORMAT(/,' Enter "1" for fast, or "0" for slow mode: ',%)
        ACCEPT 84,ISPEED
84      FORMAT(I2)
        GO TO 51

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86     TYPE 87
87     FORMAT(/, ' WHAT IS THE # FOR THE NEXT SET OF AVERAGES? ',%)
      ACCEPT 88,NSET
88     FORMAT(I4)
      GO TO 51
90     TYPE 91
91     FORMAT(/, ' MAKE HIGHER-PITCHED TONE COMMON(C) OR RARE(R)? : ',%)
      ACCEPT 92,IDHINHZ      ! Identify C or R for high-pitch tone
92     FORMAT(A1)

      IEPOCH=40*NTIX      ! Reentry point after setting parameters
      NCTONE=0             ! # of common tones presented
      NRTONE=0             ! # of rare tones presented
      NRSAMP=0             ! # of rare samples needed
      HICNT=0              ! Zero out common sample size
      LOCNT=0              ! Zero out rare sample size
      ITRIAL=0             ! Counter for # of trial

      DO 94 I=1,4
      LEVEL(I)=0           ! Estimator of waveform DC level
94     CONTINUE
      DO 96 L=1,400
      DO 95 K=1,4
      MATRIX(K,L)=0        ! Zero out single-trial matrix
      HISUM(K,L)=0         ! Zero out common summing matrix
      LOSUM(K,L)=0         ! Zero out rare summing matrix
95     CONTINUE
      YVAL(L)=0            ! Zero out display matrix
      XVAL(L)=2*L+150      ! Initialize abscissa values
96     CONTINUE
      ITIX=-1*NTIX        ! Update # ticks before clock overflow
97     TYPE 98
98     FORMAT(/, ' HOW MANY TRIALS ? (e.g., 35): ',%)
      ACCEPT 99,NTRIAL
99     FORMAT(I3)
      NRSAMP=NTRIAL-(NTRIAL*KHIPCT)/100
      PAUSE 'HIT RETURN TO GO'

      CALL DEVICE(-1,ID0,"164000)    ! Turn off all devices
      CALL IPOKE("170420,"0)        ! Turn off clock

C     *****EACH TRIAL BEGINS HERE*****
108    IF(LOCNT .GE. NRSAMP) GOTO 382 ! Already have enough R samples?
      ITRIAL=ITRIAL+1              ! Set next trial #
      RANNUH=RAN(INTGR1,INTGR2)
      CUTOFF=FLOAT(KHIPCT)/100.
      IF(RANNUH .GT. CUTOFF) GO TO 128 ! If true, R tone next
      CALL SCOPY('COMMON',TONE)
      NCTONE=NCTONE+1             ! Count # of common tones presented
      IF(IDHINHZ .EQ. 'R') GOTO 140
      GO TO 136
128    CALL SCOPY('RARE',TONE)
      NRTONE=NRTONE+1             ! Count # of rare tones presented
      IF(IDHINHZ .EQ. 'R') GOTO 136
      GO TO 140

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136      ID4="20002
      GO TO 144
140      ID4="100010
144      TYPE 148,ITRIAL,TONE
148      FORMAT(/,' Trial W',I3,' : ',4A1)

C      HERE, INITIALIZE OTHER STARTING VALUES
      JDATUM=0          ! Reset pointer for storing data
      ISYNOT=0          ! Initialize reject reason code
      IF(IEPOKI .EQ. 0) GOTO 185

      IF(ITRIAL .EQ. 1) GOTO 180
160      CALL GTIM(ETIME)
      CALL CVTTIM(ETIME,IHRS,IMIN,ISEC,ITCK)
      ITISEC=(3600*IHRS)+(60*IMIN)+(ISEC)-NOW ! Sec since last epoch
      IF(ITISEC .LE. IEPOKI) GOTO 160

180      NOW=0
185      CALL DEVICE(-1,ID1,"164000)      ! Turn on phase #1 indicator light

C      *****CLOCK WILL RUN AT 10 KHz, OVERFLOW EACH 1 MSEC*****
C      *****CLOCK INTERRUPT WILL ONLY BE USED TO UPDATE MSNEW AND ISI*****
      CALL CLOCKW(1,3,ITIX,,WATCH,"170420,"440)      ! Start clock

214      IF(JDATUM .LT. 1) GOTO 214          ! Begin CAL trigger pulse ?
      CALL DEVICE(-1,ID2,"164000)          ! CAL pulse ON
218      IF(JDATUM .LT. 25) GOTO 218          ! End CAL trigger pulse ?
      CALL DEVICE(-1,ID3,"164000)          ! CAL pulse OFF
222      IF(JDATUM .LT. 100) GOTO 222          ! Begin tone ?
      CALL DEVICE(-1,ID4,"164000)          ! Tone ON
226      IF(JDATUM .LT. 150) GOTO 226          ! End tone ?
      CALL DEVICE(-1,ID5,"164000)          ! Tone OFF
230      IF(JDATUM .LT. 400) GOTO 230          ! End trial; Stop sampling ?

      CALL IPOKE("170420,"0)      ! Turn off clock between epochs
      CALL DEVICE(-1,ID0,"164000)      ! Turn off all devices
      IF(IEPOKI .EQ. 0) GOTO 300
      CALL GTIM(ETIME)
      CALL CVTTIM(ETIME,IHRS,IMIN,ISEC,ITCK)
      ITISEC=(3600*IHRS)+(60*IMIN)+(ISEC)
      NOW=ITISEC          ! Store time at end of this trial
C      *****BEGIN INTER-TRIAL INTERVAL ACTIVITIES:*****
C      *****REVIEW DATA FOR ARTIFACTS*****
300      CALL WINDOW(MATRIX,LIMIT,LEVEL,KLIPT)
      TYPE 310,(LEVEL(N),N=1,4)
310      FORMAT(/,' DC LEVELS:',3X,I4,10X,I4,10X,I4,10X,I4)

      DO 320 I=1,4
      IF(KLIPT(I) .GT. 5) ISYNOT=4      ! More than 5 artifacts?
320      CONTINUE
      IF(ISYNOT .EQ. 4) GOTO 370

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328 IF(SCOMP('RARE',TONE) .EQ. 0) GOTO 348 ! If R trial,update LOSUM
DO 340 L=1,4
DO 332 N=1,400
HISUM(L,N)=HISUM(L,N)+MATRIX(L,N) ! Might need to convert
332 CONTINUE ! integer-to-real
340 CONTINUE
344 HICNT=HICNT+1.0
GO TO 380

348 DO 356 L=1,4
DO 352 N=1,400
LOSUM(L,N)=LOSUM(L,N)+MATRIX(L,N) ! Might need to convert
352 CONTINUE ! integer-to-real
356 CONTINUE
360 LOCNT=LOCNT+1.0
GO TO 380

370 DO 374 I=1,4
DO 372 J=1,400
MATRIX(I,J)=MATRIX(I,J) ! Keep time between tones equal
372 CONTINUE ! to when data are added to average
374 CONTINUE
TYPE 376
376 FORMAT(/,' TOO MANY ARTIFACTS; DATA NOT ADDED TO AVERAGE')

380 TYPE 381,IFIX(HICNT),IFIX(LOCNT)
381 FORMAT(' SO FAR: C =',I4,5X,'R =',I4)
IF(ISPEED .EQ. 0) GOTO 395
ICHAR=ITTINR() ! Last ASCII character entered on TT
IF(ICHAR .GT. "100") GOTO 382 ! If LETTER there, then quit speedmode
IF(LOCNT .LT. NRSAMP) GOTO 108 ! Not enough rare tone samples ?
382 TYPE 383,NCTONE,NRTONE,IFIX(HICNT),IFIX(LOCNT)
383 FORMAT(/,' # Presented : C=',I4,5X,'; R=',I4,/,
7' # Averaged : C=',I4,5X,'; R=',I4)
TYPE 385
385 FORMAT(/,' HOW MANY MORE TRIALS ? (0 if averages final) : ',%)
ACCEPT 387,NMORE
387 FORMAT(I3)
NTRIAL=NTRIAL+NMORE ! Set new max # of trials
NRSAMP=(NTRIAL*KHIPCT)/100 ! Adjust rare-sample size criterion
IF(NMORE .GT. 0) GOTO 448 ! Continue with current averages
TYPE 388
388 FORMAT(/,' PUT CURRENT AVERAGES ON FLOPPY? ',%)
ACCEPT 389,YESNO
389 FORMAT(A1)
IF(YESNO .EQ. 'Y') GOTO 670
TYPE 390
390 FORMAT(/,' BEGIN NEW SET OF AVERAGES ? ',%)
ACCEPT 391,YESNO
391 FORMAT(A1)
IF(YESNO .EQ. 'Y') GOTO 50
GO TO 448

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395 IDCH=LOOK
400 DO 404 N=1,400
    YVAL(N)=MATRIX(IDCH+1,N)+390.
404 CONTINUE
408 CALL GINIT(1)          ! Set default parameters
    CALL GEND             ! Go to transparent mode
    CALL STXOX
    CALL STOTU(7)         ! Graphic output on console
    CALL STERR(2)         ! Print error and warning messages
    TYPE 412
412 FORMAT(25(/))        ! Clear screen
    DO 416 M=1,400
    CALL POINT(XVAL(M),YVAL(M)) ! Plot values
416 CONTINUE
    CALL GEND             ! Go to transparent mode
    CALL HOLD             ! Wait for carriage return
    CALL ERASE            ! Clear screen
    CALL GEND             ! Go to transparent mode
    IF(SCOMP('DS',TASK) .EQ. 0) GOTO 464
    IF(SCOMP('DA',TASK) .EQ. 0) GOTO 476

448 TYPE 450
450 FORMAT(/, ' HIT RETURN FOR NEXT TRIAL, OR:')
    TYPE 452
452 FORMAT(' PICK FROM MENU : ??,QT,EX,WT,NU,DS,DA,LP,WD,PL,...',%)
    ACCEPT 456, TASK(1),TASK(2)
456 FORMAT(2A1)
    IF(SCOMP('??',TASK) .EQ. 0) GOTO 458 ! Display menu
    IF(SCOMP('QT',TASK) .EQ. 0) GOTO 456 ! Quit session
    IF(SCOMP('EX',TASK) .EQ. 0) GOTO 460 ! Exit session
    IF(SCOMP('WT',TASK) .EQ. 0) GOTO 382 ! Trial # status
    IF(SCOMP('NU',TASK) .EQ. 0) GOTO 50 ! Begin new set
    IF(SCOMP('DS',TASK) .EQ. 0) GOTO 464 ! Display data
    IF(SCOMP('DA',TASK) .EQ. 0) GOTO 476 ! Display average
    IF(SCOMP('LP',TASK) .EQ. 0) GOTO 600 ! List parameters
    IF(SCOMP('WD',TASK) .EQ. 0) GOTO 670 ! Write data to disk
    IF(SCOMP('PL',TASK) .EQ. 0) GOTO 648 ! Plot data
    GO TO 108

458 TYPE 460
460 FORMAT(' MENU IS AS FOLLOWS: ??---EXPLAIN MENU',/,23X,'QT---
7 QUIT SESSION (DELETE DATAFILES)',/,23X,'EX---EXIT FROM SESSION
7 (SAVE DATAFILES)',/,23X,'WT---# EACH TONE TYPE SO FAR',/,23X,
7'NU---START NEW DATA SET',/,23X,'DS---DISPLAY SINGLE TRIAL DATA',
7/,23X,'DA---DISPLAY AVERAGES',/,23X,'LP---LIST PARAMETERS',/,23X,
7'WD---WRITE DATA TO DISK',/,23X,'PL---PLOT DATA')
    GO TO 448

464 TYPE 468
468 FORMAT(' VIEW DATA FROM WHICH CHANNEL (1-4) ? ',%)
    ACCEPT 472,IDCH
472 FORMAT(I3)
    IDCH=IDCH-1
    IF(IDCH .LT. 0 .OR. IDCH .GT. 3) GOTO 448
    GO TO 400

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476 TYPE 480
480 FORMAT(' C OR R ? WHICH CHANNEL # ? (e.g. C2 or C3 or R1)',*)
ACCEPT 484,TYPE,NUMCH
484 FORMAT(A1,I1)
IF(TYPE.NE.'C'.AND.TYPE.NE.'R') GOTO 448
IF(NUMCH.LT.1 .OR. NUMCH.GT.4) GOTO 476
IF(TYPE .EQ. 'R') GOTO 492
DO 488 K=1,400
YVAL(K)=HISUM(NUMCH,K)/HICNT+390. ! Display C average
488 CONTINUE
GO TO 408
492 DO 496 K=1,400
YVAL(K)=LOSUM(NUMCH,K)/LOCNT+390. ! Display R average
496 CONTINUE
GO TO 408
600 TYPE 602,KHIPCT,LIMIT,(LOOK+1)
602 FORMAT(/,' PARAMETERS ARE AS FOLLOWS:',/, ' 1) Z OF C TONES= ',I4,
9/, ' 2) ARTIFACT LIMIT= +/- ',I6,/, ' 3) CHANNEL DISPLAYED AT END
9 OF EACH TRIAL= ',I2,/, ' 4) CAL PULSES:')
DO 606 M=1,4
TYPE 604,M,CALUV(M)
604 FORMAT(3X,'CH',I2,' ',I5,'uV')
606 CONTINUE
TYPE 608,(NTIX+40),IEPOKI,ISPEED
608 FORMAT(/,' 5) EPOCH DURATION= ',I5,' nsec',/, ' 6) INTER-EPOCH
9 INTERVAL = ',I6,' sec',/, ' 7) SPEED (0=slow;1=fast) = ',I2)
TYPE 610,IDHMHZ
610 FORMAT(/,' THE HIGHER-PITCH TONE IS THE ',A1,' TONE')
GO TO 448

648 TYPE 652
652 FORMAT(' THIS OPTION NOT YET AVAILABLE ',/)
GO TO 448

656 CALL IPOKE("170420","0") ! Shut off clock
CLOSE(UNIT=2,DISPOSE='DELETE') ! Delete datafile from DYdisk
GO TO 832

660 TYPE 664
664 FORMAT(' OK TO END THIS SESSION (Y or N): ',*)
ACCEPT 668,YESNO
668 FORMAT(A1)
IF(YESNO.NE.'Y') GOTO 448
CALL IPOKE("170420","0") ! Shut off clock
GO TO 828
C *****GENERATE LABELS FOR CURRENT DATA SET*****
670 INSTAT(1)=300 ! This was a P300 session
INSTAT(2)=MON
INSTAT(3)=IDAY
INSTAT(4)=IYEAR
DO 672 J=1,6
INSTAT(J+4)=FSPEC(J)
672 CONTINUE

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DO 673 M=1,5
  INSTAT(M+10)=IDSUBJ(M)
673  CONTINUE
  DO 675 K=1,4
    INSTAT(K+15)=CALUV(K)
675  CONTINUE
    INSTAT(20)=HICNT      ! Common tone sample size
    INSTAT(21)=LOCNT      ! Rare tone sample size
    INSTAT(22)=NTRIAL      ! Total # of trials
    INSTAT(23)=NCTONE      ! Total # common tone trials
    INSTAT(24)=NRTONE      ! Total # rare tone trials
    INSTAT(25)=NSET        ! Which # set of averages is this?
    INSTAT(26)=LIMIT       ! Artifact window size
    INSTAT(27)=NTRIAL-IFIX(HICNT+LOCNT) ! # rejects
    INSTAT(28)=NTIX        ! Tenths of msec per sample
    INSTAT(29)=IEPOKI      ! Inter-epoch interval
    INSTAT(30)=NCHS        ! # of sampled channels

D    GO TO 800
756  PRINT 760
760  FORMAT(3(/))
    CALL PUTSTR(6, IDSUBJ, ' ')
    PRINT 764, (INSTAT(I), I=1,4)
764  FORMAT(/, ' P', I3, 5X, 'Date:', I3, '/', I3, '/', I3)
    PRINT 768, INSTAT(23), INSTAT(24)
768  FORMAT(/, ' # of common tones=', I3, ' ; # of rare tones=', I3)
    PRINT 772, INSTAT(28), INSTAT(29)
772  FORMAT(/, ' Tenths of msec per sample(ms)=', I5, '/', ' ; Inter-epoch
    9 interval (sec) = ', I6)
    PRINT 776, INSTAT(26)
776  FORMAT(/, ' Artifact limit= +/-', I6, '/', '/', ' Sampled the first four
    9 channels only')
    PRINT 780, INSTAT(20), INSTAT(21)
780  FORMAT(/, ' Sample size for C average=', I3, ' ; for R average=', I3)
    PRINT 784, INSTAT(22)
784  FORMAT(/, ' Total # trials=', I3)
    PRINT 788, INSTAT(27)
788  FORMAT(/, ' Total # rejects due to artifacts=', I3)
    PRINT 796, INSTAT(16), INSTAT(17), INSTAT(18), INSTAT(19)
796  FORMAT(/, ' For each channel, CAL signal amplitude:',
    9/, 6X, 'Chan 1', 6X, 'Chan 2', 6X, 'Chan 3', 6X, 'Chan 4', /, 4(6X, I6))
    PRINT 798, INSTAT(25)
798  FORMAT(/, ' THAT WAS THE #', I4, ' SET OF AVERAGES')

C    *****WRITE DATA RECORD TO BY FILE*****
800  WRITE (2) (INSTAT(M), M=1,30)
    DO 814 I=1,4
      DO 812 J=1,400
        YVAL(J)=HISUM(I,J)/HICNT      ! Divide sum by sample size
812  CONTINUE
      WRITE (2) (YVAL(N), N=1,400)
814  CONTINUE

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      DO 824 K=1,4
      DO 820 L=1,400
      YVAL(L)=LOSUM(K,L)/LOCNT           ! Divide sum by sample size
820  CONTINUE
      WRITE (2) (YVAL(N),N=1,400)
824  CONTINUE

      TYPE 825
825  FORMAT(/,' BEGIN NEW SET OF AVERAGES ? ',0)
      ACCEPT 826,YESNO
826  FORMAT(A1)
      IF(YESNO.NE.'Y') GOTO 448
      GO TO 50                          ! Go start next set of averages
828  CLOSE(UNIT=2,DISPOSE='SAVE')      ! End this session

832  CALL EXIT
      END

      SUBROUTINE WATCH
      COMMON /BLOCK1/MATRIX,JDATUM,IBUFF
      INTEGER*2 MATRIX(4,400),IBUFF(4)
      CALL AREADP(ISTAT,IBUFF,4,-3)
      CALL EXTS12(IBUFF,4)
      JDATUM=JDATUM+1
      MATRIX(1,JDATUM)=IBUFF(1)
      MATRIX(2,JDATUM)=IBUFF(2)
      MATRIX(3,JDATUM)=IBUFF(3)
      MATRIX(4,JDATUM)=IBUFF(4)
      CALL IPOKE("170420,"133)         ! Clear overflow flag
      RETURN
      END

      SUBROUTINE WINDOW(MATRIX,LIMIT,LEVEL,KLIPT)
      INTEGER*2 MATRIX(4,400),LIMIT,LEVEL(4),KLIPT(4)
      DO 844 L=1,4
      KLIPT(L)=0                        ! Reset artifact counter
      LEVEL(L)=0                       ! Reset DC level estimator
      DO 840 M=1,400
      IF(IABS(MATRIX(L,M)).GT.LIMIT) KLIPT(L)=KLIPT(L)+1
      LEVEL(L)=LEVEL(L)+MATRIX(L,M)
840  CONTINUE
      LEVEL(L)=LEVEL(L)/400           ! Calculate mean integer value
844  CONTINUE
848  RETURN
      END

```

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C      PROGRAM PLTCNV
C      PLTCNV EMPLOYS VERTICAL CURSOR ONLY
C      Y VALUES ARE FROM ORIGINAL DATA, NOT INTERPOLATED
      EXTERNAL WATCH
      COMMON /BLOCK1/JDATUM
      VIRTUAL USAVG(8,400),DSAVG(8,400)
      REAL RLSTAT(14),USAVG,DSAVG,USCNT,DESCNT,PARAM(6)
      REAL XTABL(10),YTABL(10),XVAL(400),YVAL(400)
      LOGICAL*1 YESNO,TYPE,ICHAR,FSTRNG(14),LFILE(6),LABELX(13),LABELY(10)
      LOGICAL*1 STROFF(4),STRON(4)
      INTEGER*2 INSTAT(30),LCH(8),NMUCH
      DATA RLSTAT/14*0.0/,INSTAT/30*0/
      CALL SCOPY('Milliseconds',LABELX)           ! Abscissa label
      CALL SCOPY('Microvolts',LABELY)             ! Ordinate label
      CALL SCOPY('DY:MIXXXX.DAT',FSTRNG)          ! Initialize filename
      STROFF(1)="033
      STROFF(2)="057
      STROFF(3)="061
      STROFF(4)="144
      STRON(1)="033
      STRON(2)="057
      STRON(3)="060
      STRON(4)="144

1      TYPE 5
5      FORMAT(' Enter 4-character file code, e.g. "MI0553" : ',*)
      ACCEPT 10,(LFILE(N),N=1,6)
10     FORMAT(6A1)
      CALL INSERT(LFILE,FSTRNG,4,6)

      DO 20 L=1,400
      DO 15 K=1,8
      USAVG(K,L)=0           ! Zero out US average matrix
      DSAVG(K,L)=0           ! Zero out DS average matrix
15     CONTINUE
      YVAL(L)=0              ! Zero out display matrix
20     CONTINUE

      PARAM(1)=460.          ! XMAX
      PARAM(2)=0.            ! XMIN
      PARAM(3)=0.            ! YORDLO
      PARAM(4)=500.          ! YMAX
      PARAM(5)=-500.         ! YMIN
      PARAM(6)=-500.         ! XORDLO
      NMFLB=1

      OPEN(UNIT=2,NAME=FSTRNG,FORM='UNFORMATTED',TYPE='OLD',DISPOSE='SAVE')

      READ (2) (INSTAT(N),N=1,30),(RLSTAT(N),N=1,14)
      NMCH=INSTAT(12)
      DO 35 K=1,8
      LCH(K)=INSTAT(K+12)
35     CONTINUE

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DO 45 I=1,NCHS
READ (2) (YVAL(N),N=1,400)
DO 40 J=1,400
USAVG(LCH(I),J)=YVAL(J)
40 CONTINUE
45 CONTINUE

DO 60 K=1,NCHS
READ (2) (YVAL(N),N=1,400)
DO 55 L=1,400
DSAVG(LCH(K),L)=YVAL(L)
55 CONTINUE
60 CONTINUE

USCNT=INSTAT(21)
DSCNT=INSTAT(22)

TYPE 63
63 FORMAT(/,' Print out session info + stats ? Y or N : ',0)
ACCEPT 65,YESNO
65 FORMAT(A1)
IF(YESNO.NE.'Y') GOTO 145

PRINT 70,(INSTAT(I),I=1,4)
70 FORMAT(/,' Monkey #',I5,5X,'Date:',I3,'/',I3,'/',I3)
PRINT 75,(INSTAT(J),J=5,7)
75 FORMAT(/,' Condition #',I2,' ; Day #',I3,' ; X US trials=',I3)
PRINT 80,(INSTAT(K),K=8,10)
80 FORMAT(/,' Limited hold(ms)=',I5,/, ' Inter-stimulus interval(ms)=',
9I5,/, ' IS duration(ms)=',I6)
PRINT 85,(INSTAT(L),L=11,(INSTAT(12)+12))
85 FORMAT(/,' Artifact limit= +/-',I6,/,/, ' Sampled the following',I2,
9' channels:',8(1X,I2))
PRINT 90,INSTAT(21),INSTAT(22)
90 FORMAT(/,' Sample size for US average=',I3,' ; for DS average=',I3)
PRINT 95,INSTAT(23),INSTAT(24)
95 FORMAT(/,' Total # trials=',I3,/, ' # of US trials=',I3)
PRINT 100,(INSTAT(M),M=25,28)
100 FORMAT(/,' # aborts due to slow RT=',I3,/, ' # aborts due to premature
9 bar-pressings=',I3,/, ' # aborts due to E opting to abort=',I3,/,
9' # rejects due to artifacts=',I3)
PRINT 105,INSTAT(29),INSTAT(30)
105 FORMAT(/,' Total # bar-presses=',I5,/, ' All bar-presses during
9 ITIs=',I5)
PRINT 110,RLSTAT(1),RLSTAT(2)
110 FORMAT(/,' X US trials aborted=',F4.0,/, ' X of US trials rejected
9 due to artifacts=',F4.0)
PRINT 120,RLSTAT(3),RLSTAT(4)
120 FORMAT(/,' Mean RT(ms) for "ok" US trials=',F7.1,/,
9' Standard deviation for these RTs=',F10.4)
PRINT 125,RLSTAT(5),RLSTAT(6)
125 FORMAT(/,' For ok US trials,mean # reinforcements=',F6.3,/,
9' For ok DS trials,mean # bar-presses(in phases 6+7)=',F6.3)
PRINT 130,(RLSTAT(K),K=7,14)
130 FORMAT(/,' For artifact-rejected US trials,mean # deviant points:',
9/,2X,'Chan 1',2X,'Chan 2',2X,'Chan 3',2X,'Chan 4',2X,'Chan 5',
92X,'Chan 6',2X,'Chan 7',2X,'Chan 8',/,8(2X,F6.3))

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140 PRINT 140
140 FORMAT(5(/))
145 TYPE 150
150 FORMAT(' Display US or DS ? Channel # ? (e.g. U4 or U8 or D1) ',6)
ACCEPT 160,TYPE,NUMCH
160 FORMAT(A1,I1)
IF(TYPE.NE.'U' .AND. TYPE.NE.'D') GOTO 145
IF(NUMCH.LT.1 .OR. NUMCH.GT.8) GOTO 145
IF(TYPE .EQ. 'D') GOTO 175

DO 170 K=1,400
YVAL(K)=USAVG(NUMCH,K)
XVAL(K)=K
170 CONTINUE
GO TO 185

175 DO 180 K=1,400
YVAL(K)=DSAVG(NUMCH,K)
XVAL(K)=K
180 CONTINUE

185 TYPE 190,(PARAM(L),L=1,6)
190 FORMAT(/,' Display parameters are:',/,5X,'1)XMAX= ',F9.2,/,5X,
7'2)XMIN= ',F9.2,/,5X,'3)YGRDLO= ',F9.2,/,5X,'4)YMAX= ',F9.2,/,5X,
7'5)YMIN= ',F9.2,/,5X,'6)XGRDLO= ',F9.2)
200 TYPE 202
202 FORMAT(/,' Change display parameters ? Y or N : ',6)
ACCEPT 203,YESNO
203 FORMAT(A1)
IF(YESNO .NE. 'Y') GOTO 210
TYPE 207
207 FORMAT(/,' Enter parameter # and new value, e.g. "6,5385.7": ',6)
ACCEPT 209,NPAR,REVALU
209 FORMAT(I2,F9.2)
PARAM(NPAR)=REVALU
GO TO 185

210 DO 213 L=1,10
XTABL(L)=0
YTABL(L)=0
213 CONTINUE
XMAX=PARAM(1)
XMIN=PARAM(2)
YGRDLO=PARAM(3)
YMAX=PARAM(4)
YMIN=PARAM(5)
XGRDLO=PARAM(6)
YZERO=0.

TYPE 214
214 FORMAT(/,' "Z" to zero baseline, #s to score points, "Q" to quit')
PAUSE 'HIT RETURN'
TYPE 215
215 FORMAT(25(/))

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CALL BELL(1)
CALL ERASE
CALL GRID(LABELX,XMAX,XMIN,YGRDLO,LABELY,YMAX,YMIN,XGRDLO,MMFLG)
CALL POINT(400,XVAL,YVAL)
CALL HTEXT('X data',405.,YMAX)
CALL HTEXT('Y data',405.,-10.)
220 CALL BELL(1)
CALL VCURSR(ICHAR,XCOOR,YCOOR)
IF(ICHAR.EQ.'Q') GOTO 270
IF(ICHAR.EQ.'E') GOTO 260
IF(ICHAR.NE.'Z') GOTO 225
YZERO=YCOOR
GO TO 220
225 DECODE (1,227,ICHAR) NDATUM
227 FORMAT(I1)
IF(NDATUM.EQ.0) NDATUM=10
230 XCOOR=AIN(XCOOR+0.5) ! Reconstruct original x value
NPOS=IFIX(XCOOR)
CALL POINT(1,XVAL(NPOS),YVAL(NPOS)) ! Intensify scored point
XTABL(NDATUM)=XCOOR ! Enter x value into table
XPOS=XCOOR-3. ! Where to affix tag
CALL PLTSYM(1,XPOS,150.,ICHAR) ! Mark tag up top
VERT1=150.-10.*FLOAT(NDATUM)
VERT2=-20.-10.*FLOAT(NDATUM)
CALL HTEXT(ICHAR,405.,VERT1)
CALL FLTXT(XTABL(NDATUM),425.,VERT1) ! Display X table entry
YTABL(NDATUM)=YVAL(NPOS)-YZERO ! Non-interpolated y value
CALL HTEXT(ICHAR,405.,VERT2)
YDISP=AIN(YTABL(NDATUM)*100.)/100.
CALL FLTXT(YDISP,415.,VERT2) ! Display Y table entry
GO TO 220
240 CALL VCURSR(JCHAR,XNEW,YNEW)
DECODE (1,245,JCHAR) NDATUM
245 FORMAT(I1)
IF(NDATUM.EQ.0) NDATUM=10
CALL MVCUR(XTABL(NDATUM),YTABL(NDATUM))
XPOS=XTABL(NDATUM)-3.
YDISP=AIN(YTABL(NDATUM)*100.)/100.
VERT1=150.-10.*FLOAT(NDATUM)
VERT2=-20.-10.*FLOAT(NDATUM)
CALL PUTSTR(7,STROFF,' ') ! Data level = 1
CALL PLTSYM(1,XPOS,150.,JCHAR)
CALL FLTXT(XTABL(NDATUM),425.,VERT1)
CALL FLTXT(YDISP,415.,VERT2)
CALL PUTSTR(7,STRON,' ') ! Data level = 0
ICHAR=JCHAR
XCOOR=XNEW
YCOOR=YNEW
GO TO 230 ! Go enter new values
270 CALL ERASE
275 IF(ITTOUT('030').NE.0) GO TO 275 ! Enter transparent mode

TYPE 280
280 FORMAT(/,12X,'Datum #',BX,'X-coordinate',BX,'Y-coordinate')
DO 290 M=1,10 ! REPLACE NDATUM
TYPE 285,M,XTABL(M),YTABL(M)
285 FORMAT(12X,I3,10X,F10.3,10X,F10.3)
290 CONTINUE

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295      TYPE 295
      FORMAT(/, ' Now plot data ? : ', $)
      ACCEPT 299, YESNO
299      FORMAT(A1)
      IF(YESNO .NE. 'Y') GOTO 399
300      TYPE 305
305      FORMAT(/, ' ENTER GAIN AND OFFSET INTEGERS : ', $)
      ACCEPT 310, IGAIN, ILEVEL
310      FORMAT(I5, I5)
      JOLD=0
      JDATUM=0
      INIVAL=(YVAL(1)*IGAIN)-(ILEVEL*IGAIN)+2047
      CALL DAOUTP(1, INIVAL, "170440")
      PAUSE 'SET PLOTTER; HIT RTN'
      IPEN=1
      CALL DEVICE(-1, IPEN, "164000")
      CALL CLOCKW(1, 3, -500, WATCH, "170420, "440)
315      IF(JDATUM .LE. JOLD) GOTO 315
      JOLD=JDATUM
      IF(JDATUM .NE. 2) GOTO 317
      IPEN=0
      CALL DEVICE(-1, IPEN, "164000")
317      IF(JDATUM .GT. 400) GOTO 320
      IVAL=(YVAL(JDATUM)*IGAIN)-(ILEVEL*IGAIN)+2047
      CALL DAOUTP(1, IVAL, "170440")
      GOTO 315
320      CALL IPOKE("170420, "0)          ! Turn off clock
      TYPE 325
325      FORMAT(/, ' REPLOT DATA ? : ', $)
      ACCEPT 330, YESNO
330      FORMAT(A1)
      IF(YESNO .EQ. 'Y') GOTO 300

399      TYPE 400
400      FORMAT(/, ' View more average data (Y or N) ? ', $)
      ACCEPT 410, YESNO
410      FORMAT(A1)
      IF(YESNO .EQ. 'Y') GOTO 145

      CLOSE(UNIT=2, DISPOSE='SAVE')
      TYPE 430
430      FORMAT(/, ' OPEN NEW DATAFILE ? : ', $)
      ACCEPT 435, YESNO
435      FORMAT(A1)
      IF(YESNO .EQ. 'Y') GOTO 1

      CALL EXIT
      END

      SUBROUTINE WATCH
      COMMON /BLOCK1/JDATUM
      JDATUM=JDATUM+1
      CALL IPOKE("170420, "133)
      RETURN
      END

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**END**

**FILMED**

**1-84**

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